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For: MOLECULAR INTERACTIONS IN HEMATOPOIETIC CELLS

- [X] This application claims priority from each of the following Application Nos./filing dates:
60/196,460 filed April 11, 2000, 60/196,528 filed April 11, 2000, 60/196,527 filed April 11, 2000, 60/196,267 filed April 11, 2000, 60/182,296 filed February 14, 2000, 60/176,195 filed January 14, 2000, 60/170,453 filed December 13, 1999, 60/162,498 filed October 29, 1999, 60/160,860 filed October 21, 1999, 60/134,118 filed May 14, 1999, 60/134,117 filed May 14, 1999, 60/134,114 filed May 14, 1999, Unassigned filed October 13, 2000, 09/570,118 filed May 12, 2000, 09/570,34 filed May 12, 2000, 09/569,525 filed May 12, 2000, 09/547,276 filed April 11, 2000
the disclosure(s) of which is (are) incorporated by reference.

Enclosed are:

- [X] 181 page(s) of specification
[X] 5 page(s) of claims
[X] 1 page of Abstract
[X] 19 sheet(s) of ☐ formal ☒ informal drawing(s).
[X] 1 Title page.

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[X] Copies of Petitions (3) to extend time to respond in parent applications App. Nos. 09/570,118; 09/570,364; 09/569,525.

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BASIC FEE		
TOTAL CLAIMS	17 - 20	= *0
INDEP. CLAIMS	3 - 3	= *0
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PATENT APPLICATION
MOLECULAR INTERACTIONS IN HEMATOPOIETIC CELLS

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MOLECULAR INTERACTIONS IN HEMATOPOIETIC CELLS

REFERENCE TO RELATED APPLICATIONS

5 This application claims priority to U.S. patent application No. _____ (filed November 10, 2000, attorney docket no. 020054-001120US), U.S. patent application No. _____ (filed October 13, 2000, attorney docket no. 020054-001110US), U.S. patent applications serial nos. 09/570118, 09/570364, 09/569525 (all filed May 12, 2000), 60/196,460, 60/196,528, and 60/196,527, 60/196,267, 09/547,276 (all filed April 11, 2000), serial no. 10 60/182,296 (filed February 14, 2000); serial no. 60/176,195 (filed January 14, 2000); serial no. 60/170,453 (filed December 13, 1999); serial no. 60/162,498 (filed October 29, 1999); serial no. 60/160,860 (filed October 21, 1999); and serial nos. 60/134,118; 60/134,117; and 60/134,114 (all filed May 14, 1999); the disclosures of each of which are incorporated herein in their entirety.

1. FIELD OF THE INVENTION

15 The present invention relates to peptides and peptide analogues, and methods for using such compositions to regulate activities of cells of the hematopoietic system. In one aspect, the invention provides methods of modulating metabolism (e.g., activation) of hematopoietic cells (e.g., T cells and B cells) by antagonizing an interaction between a PDZ domain containing protein and a protein that binds a PDZ domain. In one aspect, it relates to 20 fusion peptides containing an amino acid sequence corresponding to the carboxyl terminus of a surface receptor expressed by a hematopoietic cell and a transmembrane transporter sequence; such fusion peptides are useful in regulating hematopoietic cells by inhibiting cell activation.

2. BACKGROUND OF THE INVENTION

25 PDZ domains of proteins are named after three prototypical proteins: PSD95, Drosophila large disc protein and Zonula Occludin 1 protein (Gomperts et al., 1996, *Cell* 84:659-662). PDZ domain-containing proteins are involved in synapse formation by organizing transmembrane neurotransmitter receptors through intracellular interactions. PDZ domains contain the signature sequence GLGF (SEQ ID NO: 402). In the nervous system, typical PDZ domain-containing proteins contain three PDZ domains, one SH3 domain and one guanylate 30 kinase domain. Examples of intracellular PDZ domain-containing proteins include LIN-2, LIN-7 and LIN-10 at the pre-synapse, and PSD95 at the post-synapse.

PDZ domains have been shown to bind the carboxyl termini of transmembrane proteins in neuronal cells. Songyang et al. reported that proteins capable of binding PDZ domains contain a carboxyl terminal motif sequence of E-S/T-X-V/I (Songyang et al., 1997, *Science* 275:73). X-ray crystallography studies have revealed the contact points between the motif sequence and PDZ domains (Doyle et al., 1996, *Cell* 88:1067-1076). While the interaction between PDZ domains and ion channels in neurons have been studied extensively, such interactions have had limited studies in other biological systems, especially the hematopoietic system.

The hematopoietic system is composed of different cell types that perform distinct functions. Many of its diverse functions require coordinated movement of cell surface receptors including ion channels, adhesion surface molecules to coordinate cell-cell interaction, and cytokine receptors. Despite their diverse functional activities, all hematopoietic cells are believed to develop from a multipotent bone marrow hematopoietic stem cell. Such stem cell has been shown to express a surface marker termed CD34. During differentiation, the stem cell gives rise to progenitor cells in each of several specific hematopoietic cell lineages. The progenitor cells then undergo a series of morphological and functional changes to produce mature functionally committed hematopoietic cells.

Among the functions performed by hematopoietic cells, certain cell types are involved exclusively in immunity. For example, lymphocytes, which include T cells, B cells and natural killer (NK) cells, are effectors in immune responses. Monocytes and granulocytes (i.e., neutrophils, basophils and eosinophils) play a role in non-specific forms of defense. Lymphocytes, monocytes and granulocytes are collectively referred to as white blood cells or leukocytes. On the other hand, other hematopoietic cells perform functions that are unrelated to the immune system. For example, erythrocytes are involved in gas transport, and cells of the thrombocytic series are involved in blood clotting.

T cells and B cells recognize antigens and generate an immune response. T cells recognize antigens by heterodimeric surface receptors termed the T cell receptor (TCR). The TCR is associated with a series of polypeptides collectively referred to as CD3 complex. B cells recognize antigens by surface immunoglobulins (Ig), which are also secretory molecules. In addition, a large number of co-stimulatory surface receptors have been identified in T cells and B cells, which augment cellular activation during antigen-induced activation.

In addition to the T cell antigen receptor/CD3 complex (TCR/CD3), other molecules expressed by T cells which mediate an activation signal, include but are not limited to, CD2, CD4, CD5, CD6, CD8, CD18, CD27, CD28, CD43, CD45, CD152 (CTLA-4), CD154, MHC class I, MHC class II, CDw137 (4-1BB), CDw150, and the like (Barclay et al., The

Leucocyte Antigen Facts Book, 1997, Second edition, Academic Press; Leucocyte Typing, 1984, Bernard et al. (eds.), Springer-Verlag; Leukocyte Typing II, 1986, Reinherz et al. (eds.), Springer-Verlag; Leukocyte Typing III, 1987, McMichael (ed.), Oxford University Press; Leukocyte Typing IV, 1989, Knapp et al. (eds.), Oxford University Press; CD Antigens, 1996, VI Internat. Workshop and Conference on Human Leukocyte Differentiation Antigens. <http://www.ncbi.nlm.nih.gov/prow>); all incorporated by reference herein. Cell surface antigens that work together with TCR/CD3 are often referred to as co-receptors in the art.

Specific antibodies have been generated against all of the aforementioned T cell surface antigens. Other molecules that bind to the aforementioned T cell surface receptors include antigen-binding antibody derivatives such as variable domains, peptides, superantigens, and their natural ligands such as CD58 (LFA-3) for CD2, HIV gp120 for CD4, CD27L for CD27, CD80 or CD86 for CD28 or CD152, ICAM1, ICAM2 and ICAM3 for CD11a/CD18, 4-1BBL for CDw137.

Activation molecules expressed by B cells, include but are not limited to, surface Ig, CD18, CD19, CD20, CD21, CD22, CD23, CD40, CD45, CD80, CD86 and ICAM1. Similarly, natural ligands of these molecules and antibodies directed to them as well as antibody derivatives may be used to deliver an activation signal to B cells.

However, prior to the present invention, it was not known that signal transduction following stimulation of any leukocyte receptor was mediated by receptor interactions with PDZ domain-containing proteins. Therefore, it was not even contemplated in the art that an interference of leukocyte surface receptor/PDZ domain interactions could regulate leukocyte activation.

3. SUMMARY OF THE INVENTION

In one aspect, the invention provides a method of modulating a biological function of a cell, e.g., an endothelial cell or hematopoietic cell (such as a leukocyte, e.g., T cell or B cell), by introducing into the cell an antagonist that inhibits binding of a PDZ protein and a PL protein in the cell, or a agonist that enhances binding of a PDZ protein and a PL protein in the cell. In various embodiments the PL protein is an adhesion protein, an adaptor protein, or an intracellular protein. In embodiments it is CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD4, CD97, CD3n, DOCK2, CD34, FcεRIb, or FasLigand. In an embodiment the PL protein is characterized by a carboxy-terminal amino acid motif that is X-S-X-A, X-A-D/E-V, X-V/I/L-X*-V, or X-S/T-X-F (where X is any amino acid and X* is any non-aromatic amino acid). In embodiments, the PL protein is expressed by T lymphocytes or B lymphocytes. In some embodiments of this method, the PDZ protein is CASK, MPP1, DLG1, PSD95, NεDLG, SYN1a, TAX43, LDP, LIM, LIMK, AF6, PTN-4, prIL16, 41.8, RGS12,

DVL1, TAX 40, TIAM1, MINT1, K303, TAX2, or KIAA561.

In some embodiments, the cell is a leukocyte and the biological function is cell activation, cell proliferation, maintenance of cell structure, cell metabolic activity, or cytokine production. In some embodiments, the method further includes detecting a change in leukocyte activation.

In preferred embodiments, the antagonist is an agent that inhibits the binding of a PL peptide to a PDZ domain polypeptide in an "A" assay, in a "G" assay, or in both an A assay and a G assay. The antagonist can be a polypeptide, such as a polypeptide having at the carboxyterminus at least two residues that are the same as the carboxy-terminal two residues of a PL protein, such as a PL protein is expressed in a hematopoietic or endothelial cell, and/or that is an adhesion protein, an adaptor protein, or an intracellular protein. In an embodiment, at least the carboxy-terminal four residues of the polypeptide are the same as the carboxy-terminal four residues of the PL protein. In an embodiment, the PL protein has a carboxy-terminal amino acid motif selected from X-S-X-A, X-A-D/E-V, X-V/I/L-X*-V, or X-S/T-X-F, where X is any amino acid and X* is any non-aromatic amino acid. In embodiment, the PL protein is CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD97, CD3n, DOCK2, CD34, FcεRIb, or FasLigand.

In a related aspect, the antagonist is a peptide mimetic of a PL inhibitor sequence peptide. In another related aspect the antagonist is a fusion polypeptide having a PL sequence and transmembrane transporter amino acid sequence (such as HIV tat, Drosophila antenapedia, herpes simplex virus VP22 or anti-DNA CDR 2 and 3).

In another aspect, the invention provides a method of determining whether a test compound is an inhibitor of binding between a PDZ protein and a PL protein by contacting a PDZ domain polypeptide having a sequence from the PDZ protein, and a PL peptide under conditions in which they form a complex, in the presence and in the absence of a test compound, and detecting the formation of the complex in the presence and absence of the test compound, where less complex formation in the presence of the test compound than in the absence of the compound indicates that the test compound is an inhibitor of a PDZ protein -PL protein binding. In embodiments the PL peptide has a sequence that includes the a C-terminal sequence of a PL protein, such as CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD97, CD3n, DOCK2, CD34, FcεRIb, or FasLigand. In some embodiments, the PDZ domain polypeptide is a fusion polypeptide.

In a related aspect, the invention provides a method of determining whether a test compound is an agonist of binding between a PDZ protein and a PL protein by contacting a PDZ domain polypeptide, and a PL peptide under conditions in which they form a complex, in the

presence and in the absence of a test compound, and detecting the formation of the complex in the presence and absence of the test compound, where more complex formation in the presence of the test compound than in the absence of the compound indicates that the test compound is an agonist of a PDZ protein-PL protein binding.

5 The invention further provides an inhibitor of binding of a PDZ protein and a PL protein. In an embodiment, the inhibitor is characterized in that it reduces binding of a peptide selected from the group consisting of a PL peptide selected from the group consisting of CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD97, CD3n, DOCK2, CD34, FcεRIb, and FasLigand and a PDZ domain polypeptide. In
10 various embodiments, the inhibitor is a peptide comprising a sequence that is from 3 to about 20 residues of a C-terminal sequence of a PL protein selected from CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD97, CD3n, DOCK2, CD34, FcεRIb, and FasLigand; a peptide having a motif X-S-X-A, X-A-D/E-V, X-V/I/L-X*-V, or X-S/T-X-F, (where X is any amino acid and X* is any non-aromatic amino acid); a peptide
15 mimetic; or a small organic molecule. The invention also provides a pharmaceutical composition containing the inhibitor.

The invention also provides a method for treating a disease characterized by leukocyte activation by administering a therapeutically effective amount of an inhibitor of a PL-PDZ interaction. In embodiments, the disease is characterized by an inflammatory or humoral
20 immune response or is an autoimmune disease. The invention further provides a method of reducing inflammation in a subject by administering an agent that inhibits binding of a PDZ protein and a PL protein, where the PL protein is an adhesion protein, an adaptor protein, or an intracellular protein.

The invention also provides use of an inhibitor of the binding of a PDZ protein
25 and a PL protein to inhibit leukocyte activation or to treat a disease mediated by hematopoietic cells, such as a disease is characterized by an inflammatory or humoral immune response. The invention also provides use of an inhibitor of the binding of a PDZ protein and a PL protein in the preparation of a medicament for treatment of a disease mediated by hematopoietic cells.

The invention also provides a method of modulating a biological function of a
30 hematopoietic cell, comprising introducing into the cell an antagonist that inhibits binding of a PDZ protein and a PL protein in the cell as deduced from Table 2, for example, where the PL protein is DNAM-1 and the PDZ protein is MPP1, MPP2, DLG1, NeDLG, PSD95, LIM, AF6, 41.8 or RGS12, the PL protein is LPAP and the PDZ protein is DLG1 or MINT1, or the PL protein is DNAM-1 and the PDZ protein is PSD95 or MPP2.

The present invention also relates to peptides and peptide analogues that bind PDZ domains in hematopoietic cells. In particular, it relates to fusion peptides and peptide analogues containing a hematopoietic cell surface receptor carboxyl terminal sequence and a transmembrane transporter sequence which facilitates entry of the peptides into a target cell. The invention also relates to methods of using such compositions in inhibiting leukocyte activation as measured by cytokine production, cell proliferation, apoptosis and/or cytotoxicity.

It is an object of the invention to administer a therapeutically effective amount of the aforementioned fusion peptides, peptide analogues, small molecules and other mediators of PDZ-PL interactions as pharmaceutical compositions, e.g., to a subject to inhibit undesirable cell-mediated (e.g., leukocyte-mediated) events.

It is also an object of the invention to administer a therapeutically effective amount of the aforementioned fusion peptides, peptide analogues, small molecules and other mediators of PDZ-PL interactions as pharmaceutical compositions to a subject to treat an autoimmune disorder or to prevent transplantation rejection of a solid organ transplant.

In one aspect, the invention provides a method of determining the apparent affinity (K_d) of binding between a PDZ domain and a ligand by (a) immobilizing a polypeptide comprising the PDZ domain and at least one non-PDZ domain on a surface; (b) contacting the immobilized polypeptide with a plurality of different concentrations of the ligand; (c) determining the amount of binding of the ligand to the immobilized polypeptide at each of the concentrations of ligand; (d) calculating the apparent affinity of the binding from the binding determined in (c). In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-PDZ domain. In an embodiment, the polypeptide comprising the PDZ domain is a fusion protein, for example a GST-PDZ domain fusion protein.

In one aspect, the invention provides a method of determining the K_i of an inhibitor or suspected inhibitor of binding between a PDZ domain and a ligand, by (a) immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface; (b) contacting the immobilized polypeptide with a plurality of different mixtures of the ligand and inhibitor, wherein the different mixtures comprise a fixed amount of ligand, at least a portion of which is detectably labeled, and different concentrations of the inhibitor; (c) determining the amount of ligand bound at the different concentrations of inhibitor; (d) calculating the K_i of the inhibitor from the binding determined in (c). In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-PDZ domain. In an embodiment, the fixed amount of ligand is between about 0.01 K_d and about 2 K_d .

In another aspect, the invention provides a method of identifying an agent that enhances the binding of a PDZ domain to a ligand, by immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface; (b) contacting the immobilized polypeptide with the ligand in the presence of a test agent and determining the amount of ligand bound; and,
5 (c) comparing the amount of ligand bound in the presence of the test agent with the amount of ligand bound by the polypeptide in the absence of the test agent, wherein at least two-fold greater binding in the presence of the test agent compared to the absence of the test agent indicates that the test agent is an agent that enhances the binding of the PDZ domain to the ligand. In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized
10 immunoglobulin that binds the non-PDZ domain.

In another aspect, the invention provides a method of determining the potency (K_{enhancer}) of an enhancer of binding between a PDZ domain and a ligand, by (a) immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface; (b) contacting the immobilized polypeptide with a plurality of different mixtures of the ligand and enhancer,
15 wherein the different mixtures comprise a fixed amount of ligand, at least a portion of which is detectably labeled, and different concentrations of the enhancer; (c) determining the amount of ligand bound at the different concentrations of enhancer; (d) calculating the potency (K_{enhancer}) of the enhancer from the binding determined in (c). In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-
20 PDZ domain. In an embodiment, the fixed amount of ligand is between about 0.01 Kd and about 0.5 Kd.

In another aspect, the invention provides a method of identifying a high specificity interaction between a particular PDZ domain and a ligand known or suspected of binding at least one PDZ domain, by (a) providing a plurality of different immobilized polypeptides, each of said
25 polypeptides comprising a PDZ domain and a non-PDZ domain; (b) determining the affinity of the ligand for each of said polypeptides; (c) comparing the affinity of binding of the ligand to each of said polypeptides. An interaction between the ligand and a particular PDZ domain is deemed to have high specificity when the ligand binds an immobilized polypeptide comprising the particular PDZ domain with at least 2-fold higher affinity than to immobilized polypeptides
30 not comprising the particular PDZ domain in (a). In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-PDZ domain.

In another aspect, the invention provides a method for determining the PDZ-PL inhibition profile of a compound by (a) providing (i) a plurality of different immobilized polypeptides, each of said polypeptides comprising a PDZ domain and a non-PDZ domain; (ii) a plurality of corresponding ligands, wherein each ligand binds at least one PDZ domain in (i);
5 (b) contacting each of said immobilized polypeptides in (i) with a corresponding ligand in (ii) in the presence and absence of a test compound; (c) determining for each polypeptide-ligand pair in (b) whether the test compound inhibits binding between the immobilized polypeptide and the corresponding ligand thereby determining the PDZ-PL inhibition profile of the test compound.

In another aspect, the invention provides an array comprising a plurality of
10 different immobilized polypeptides, each of said polypeptides comprising a PDZ domain and a non-PDZ domain. In an embodiment, the array is situated in a plastic multiwell plate. In an embodiment, the array has at least 12 different polypeptides comprising at least 12 different PDZ domains, for example, at least 12 different PDZ domains are from PDZs expressed in lymphocytes. In an embodiment, the PDZs are selected from those listed in Table 2 or 6.

In an aspect, the invention provides an assay device comprising a plurality of
15 different immobilized PDZ-containing proteins organized in an array. In one embodiment, the device has at least 25 different PDZ-containing proteins.

In a further aspect, the invention provides a method for identifying an interaction between a PDZ domain and a PL by contacting a PL to a plurality of PDZ containing polypeptides and detecting binding of at least one PL to a PDZ. In an embodiment, the
20 contacting occurs on an assay device comprising a plurality of different immobilized PDZ-containing proteins organized in an array. In one embodiment, the device has at least 25 different PDZ-containing proteins. In embodiments, an interaction between a PDZ and more than one PL, or between a PL and more than one PDZ, is detected.

In a related aspect, the invention provides method for identifying a modulator of
25 an interaction between a PDZ and a PL by conducting any of the aforementioned assays in the presence and absence of a test compound and detecting a difference in at least one PDZ-PL interaction in the presence and absence of the test compound. In embodiments, the the modulator is an enhancer of the interaction. In other embodiments, the modulator is an inhibitor of the
30 interaction.

In an embodiment, any of the aforementioned methods or devices (as further described herein) comprising a plurality of PDZ-domain containing polypeptides (e.g., a PDZ domain fusion protein) comprises at least one, usually at least 2, typically at least 5 and often at

least 10 different PDZ-containing polypeptides comprising PDZ sequences from proteins selected from: MPP1 (p55), K303, K807, DLG1, PSD95, NeDLG, TAX IP43, LDP, LIM, K545, TIP1, PTN-4, CBP, AF6, PDZK1, DLG5, Syntenin, WWP3, K561.

In an aspect, the invention provides a method of modulating a biological function of an endothelial cell or hematopoietic cell (e.g., a leukocyte such as a T cell or a B cell), comprising introducing into the cell an agent that inhibits binding of a PDZ protein and a PL protein in the cell, wherein any of the following (I)-(I) apply:

(I) the PL is CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, CD44, CD38, CD3 η , LPAP, CD46, CDw128B, DOCK2, PAG, CD34, or BLR-1;

(II) the PDZ is MPP1, K303, K807, DLG1, PSD95, NeDLG, IP43, LDP, LIM, K545, TIP1, PTN-4, CBP, AF6, PDZK1, DLG5, Syntenin, WWP3, or K561;

(III) the PDZ protein is MPP1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y/I-X-V; the PDZ protein is LIMK1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V; the PDZ protein is K303 and the PL protein has a carboxy-terminal amino acid motif X-S-X-V; the PDZ protein is K807 and the PL protein has a carboxy-terminal amino acid motif X1-S/T-X2-V/I/L/F; the PDZ protein is DLG1, PSD95, or NeDLG and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y/A/E-X-V/I/L; the PDZ protein is SNTa1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-D/Y-V/I/L; the PDZ protein is DVL1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V; the PDZ protein is LDP and the PL protein has a carboxy-terminal amino acid motif X-A/S-X2-V/I; the PDZ protein is LIM and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-A/V; the PDZ protein is K561 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V/I/L/F; the PDZ protein is K545 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y-M-A/S/V; the PDZ protein is TAX-IP2 and the PL protein has a carboxy-terminal amino acid motif X-S-D/E-V; the PDZ protein is MPP2 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-A/V/I; the PDZ protein is TIP-1 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-V/I/L; the PDZ protein is PTN-4 and the PL protein has a carboxy-terminal amino acid motif X1-S/T-X-V/F; the PDZ protein is prIL16 and the PL protein has a carboxy-terminal amino acid motif D/E/K/R-V/I/L/F/Y-X-V; the PDZ protein is CBP and the PL protein has a carboxy-terminal amino acid motif X-S/T-F/Y-V; the PDZ protein is protein 41 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y/F-X-A/V/I/L; the PDZ protein is AF6 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y-F/Y-V/I/L; the PDZ protein is RGS12 and the

PL protein has a carboxy-terminal amino acid motif X1-S/T/Y-X-V/F; the PDZ protein is PDZK1 and the PL protein has a carboxy-terminal amino acid motif X-T-X-F; the PDZ protein is DLG5 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X-V; the PDZ protein is Synt and the PL protein has a carboxy-terminal amino acid motif X1-V/I/L-X2-V; the PDZ protein is WWP3 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-V; or the PDZ protein is TAX-IP40 and the PL protein has a carboxy-terminal amino acid motif X-Y-X-V; where X is any amino acid, X1 is any amino acid, X2 is any amino acid;

(IV) the agent is a peptide comprising a sequence of at least the carboxy-terminal two or three residues of the PL protein;

(V) the agent is a small molecule or peptide mimetic of the carboxy-terminus of the PL protein;

In an aspect the invention provides a method for determining whether a test compound is an inhibitor of binding between a PDZ protein and a PL protein by contacting a PDZ domain polypeptide having a sequence from the PDZ protein, and a PL peptide, wherein the PL peptide comprises a C-terminal sequence of a PL protein under conditions in which they form a complex, where the contacting is carried out in the presence and in the absence of a test compound, and detecting the formation of the complex in the presence and absence of the test compound. In embodiments, the PL protein is CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, TAX, CD44, CD38, CD3 η , LPAP, CD46, CDw128B, DOCK2, PAG, CD34, or BLR-1 and less complex formation in the presence of the test compound than in the absence of the compound indicates that the test compound is an inhibitor of a PDZ protein-PL protein binding. The invention also contemplates the inhibitor identified by this method. In embodiments, the inhibitor is (a) a peptide comprising a sequence that is from 3 to about 20 residues of a C-terminal sequence of CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, TAX, CD44, CD38, CD3 η , LPAP, CD46, CDw128B, DOCK2, PAG, CD34, or BLR-1; (b) a peptide mimetic of such a peptide; or (c) a small organic molecule with a molecular weight less than 1 kD. The invention further contemplates a pharmaceutical composition containing the inhibitor, as well as a method for treating a disease characterized by leukocyte activation by administering a therapeutically effective amount of the inhibitor. In embodiments, the disease is characterized by an inflammatory or humoral immune response, e.g., an autoimmune disease.

In an aspect, the invention provides a method of modulating a biological function in a cell (e.g., a hematopoietic cell) by introducing into the cell an antagonist that inhibits binding of a PDZ protein and a PL protein in the cell, wherein, the PDZ protein is MPP1 (p55) and the PL is Spectrin β ; the PDZ protein is K303 and the PL is Spectrin β ; the PDZ protein is K807 and

the PL VCAM1, Spectrin β , KV1.3, Neuroligin 3, CD38, CD3 η , LPAP, CD46 (form 1), CDw128B, DOCK2, PAG, CD34, or BLR-1; the PDZ protein is DLG1 and the PL is Spectrin; the PDZ protein is PSD95 and the PL is Spectrin β , CD34, or CD38; the PDZ protein is NeDLG and the PL is Spectrin β or CD38; the PDZ protein is TAX IP43 and the PL is Spectrin β or CD38; the PDZ protein is LDP and the PL is CD38; the PDZ protein is LIM and the PL is CD105; the PDZ protein is K545 and the PL is CD105; the PDZ protein is TIP1 and the PL is CD95, KV1.3, CD3 η , LPAP; the PDZ protein is PTN-4 and the PL is Spectrin β ; the PDZ protein is CBP and the PL is Spectrin β ; the PDZ protein is AF6 and the PL is Spectrin β ; the PDZ protein is PDZK1 and the PL is BLR-1; the PDZ protein is DLG5 and the PL is Spectrin; the PDZ protein is Syntenin and the PL is CD44; the PDZ protein is WWP3 and the PL is VCAM1, Spectrin β , DNAM1, Neuroligin 3; the PDZ protein is K561 and the PL is BLR-1.

The invention also provides the use of an inhibitor of the binding of a PDZ protein and a PL protein described herein or identified according to a method of the invention to inhibit leukocyte activation, or for preparation of a medicament for treatment of a disease mediated by a PDZ-PL interaction, e.g., in hematopoietic cells or in viral infection.

The PDZ and PL proteins referred to herein are known in the art and are described herein, e.g., at Tables 3, 4 and 7. For example, CD105 is described at GenBank accession no. X72012; VCAM1 is described at GenBank accession no. M73255; CD95 is described at GenBank accession no. M67454; Spectrin β is described at GenBank accession no. NM000347; KV1.3 is described at GenBank accession no. AAC31761; DNAM1 is described at GenBank accession no. U56102; Neuroligin 3 is described at GenBank accession no. NM018977; TAX is described at GenBank accession no. AB038239; CD44 is described at GenBank accession no. M69215; CD38 is described at GenBank accession no. NM004334; CD3 η is described at GenBank accession no. M33158; LPAP is described at GenBank accession no. X81422; CD46 is described at GenBank accession no. M58050; CDw128B is described at GenBank accession no. M73969; DOCK2 is described at GenBank accession no. BAA13200; PAG is described at GenBank accession no. NM018440; CD34 is described at GenBank accession no. M81104; BLR-1 is described at GenBank accession no. S56162; CD4 is described at GenBank accession no. M12807; CD6 is described at GenBank accession no. X60992; CD49E (4) is described at GenBank accession no. X06256; CD49F is described at GenBank accession no. X53586; CD97 is described at GenBank accession no. X84700; CD98 is described at GenBank accession no. J02939; CD138 is described at GenBank accession no. J05392; CD148 is described at GenBank

accession no. D37781; CD166 is described at GenBank accession no. L38608; CDw137 (4-1BB) is described at GenBank accession no. NM001561; FasL is described at GenBank accession no. U11821; FcεRIβ is described at GenBank accession no. D10583; Galectin3 is described at GenBank accession no. J02921; CD114 is described at GenBank accession no. NM000760; CDW125 (IL5R) is described at GenBank accession no. X62156; CDW128A (IL8RA) is described at GenBank accession no. M68932; Mannose Receptor is described at GenBank accession no. NM002438; NMDA is described at GenBank accession no. NP000824; Glycophorin C is described at GenBank accession no. AAA52574; Neurexin is described at GenBank accession no. AB011150; Syndecan-2 is described at GenBank accession no. A33880; CC CKR-1R is described at GenBank accession no. L09230; CC CKR-2 is described at GenBank accession no. U03882; CC CKR-3 is described at GenBank accession no. HSU28694; CC CKR-4 is described at GenBank accession no. X85740; Volt. Gated Ca²⁺ is described at GenBank accession no. Q00975; CD83 is described at GenBank accession no. Z11697; CD62E is described at GenBank accession no. M30640; CD5 is described at GenBank accession no. X04391; and CD148 is described at GenBank accession no. D37781; BLR-1/CXCR5 NM001716.

4. BRIEF DESCRIPTION OF THE DRAWINGS

FIGURES 1A-1D show the results of exemplary assays in which the binding of biotinylated peptides having a sequence of the carboxyl-terminus ("c-terminus") of various leukocyte proteins to PDZ domains (i.e., GST-PDZ domain fusion proteins) was determined using the "G" assay described *infra*. The PDZ domains are: PSD95 (Fig. 1A); NeDLG (Fig. 1B); DLG1 (Fig 1C); and 41.8 (Fig. 1D). These and other PDZ domain fusion proteins are described *infra* (e.g., TABLE 2). In the figure, peptides 1-31 refer to the biotinylated PL peptides used in the assay, and are identified in the Key, *infra*. "Peptide IDs" are defined in TABLE 3. Key:

#	Test Protein	Peptide IDs
1	Clasp-2	AA2L
2	FceRI β	AA25L
3	CDW128B	AA29.2
4	KV1.3	AA33L
5	Neurexin	AA38L
6	DOCK2	AA40L
7	CC CKR-1R	AA41L
8	CC CKR-2	AA42L
9	CC CKR-4	AA44L
10	BLR-1	AA45L
11	CD49E	AA11L
12	CD97	AA14L
13	VCAM1	AA17L
14	CD138	AA18L
15	DNAM-1	AA22L

16	CDW128A	AA29.1L
17	CC CKR-3	AA43L
18	Clasp-1	AA1L-R
19	CD46 (Form 1)	AA10L
20	CD95	AA13L
21	CDW125	AA28L
22	CD83	AA47L
23	CD62E	AA48L
24	CD3n	AA4L
25	Clasp-4	AA3L-V
26	CD44	AA9L
27	CD166	AA20L
28	CD62E	AA48L
29	CD5	AA49L
30	CD148	AA55L
31	DOCK2	AA40L

FIGURES 2A and 2B show the Apparent Affinity Determination for PDZ–Ligand Interactions. Varying concentrations of biotinylated CLASP-2 (Fig. 2A; **TABLE 4**) or Fas (Fig. 2B; **TABLE 4**) C-terminal peptides were reacted with immobilized (plate bound) GST polypeptide or GST-PDZ fusion proteins (GST-DLG1, GST-NeDLG, and GST-PSD95). The binding to GST alone (< 0.2 OD units) was subtracted from the binding to the fusion proteins to obtain the signal at each peptide concentration. This signal was then normalized by dividing the signal at each peptide concentration by the maximum signal observed for each peptide-PDZ pair (i.e. the signal obtained at 30 μ M Clasp 2 peptide or 100 μ M Fas peptide; 0.4 – 1.0 OD units for Clasp 2 and 1.2 – 2.0 OD units for Fas). The normalized signals were then plotted and fit to a saturation binding curve, yielding an apparent affinity of 21 μ M for DLG1-Clasp 2 interaction, 7.5 μ M for NeDLG-Clasp 2 interaction, 45 μ M for PSD95-Clasp 2 interaction, 54 μ M for DLG1-Fas interaction, 54 μ M for NeDLG-Fas interaction, and 85 μ M for PSD95-Fas interaction. Data are means of duplicate data points, with standard errors between duplicate data points $< 20\%$.

FIGURES 3A-3F show inhibition of PDZ – PL peptide interactions. A fixed concentration of biotinylated C-terminal peptide having a sequence based on the C-terminal sequence of a cell surface receptor protein (Clasp 2, CD46, Fas, and KV1.3; see **TABLE 4**) was bound to immobilized GST polypeptide or the GST-fusion protein indicated at the top left of each frame, in the presence or absence of the competitor peptides indicated in the legend of each frame and the level of inhibition determined. Fig. 3A- DLG1; Fig. 3B- PSD95; Fig. 3C NeDLG; Fig. 3D - DLG1, Fig. 3E, PSD95; Fig. F - 41.8. In Fig. 3A-B the competitor peptides

are present at 100 uM; in Figs. 3C-F the competitor is present at the indicated concentration.

FIGURES 4A and 4B shows the results of introduction of a Tat-CD3 fusion peptide on T cell activation. Antigen-specific T cell activation was measured by cytokine production. Fusion peptides containing tat and a T cell surface molecule carboxyl terminus inhibited γ -interferon (IFN) production by a T cell line in response to myelin basic protein (MBP) stimulation. The level of inhibition was determined by first subtracting the binding of the labeled peptide to GST alone from the binding to the fusion protein and dividing by the signal in the absence of competitor peptide.

FIGURES 5A and 5B show TIP1-RFP overexpression enhances anti-CD95 induced apoptosis in Jurkat T cells. Jurkat E6 T cells were transfected with either DsRED (RFP), TIP1-RFP, or PAR6(N-P)-RFP. 24 hours post- transfection, cells were treated for 2 hours with anti-CD95 and then incubated with annexin V-FITC and analyzed using flow cytometry. Fig. 5A shows the results of FACS analysis. Fig. 5B shows a 30% increase in apoptosis of RFP-TIP1 positive compared to RFP negative apoptotic (annexin V positive) cells.

FIGURE 6 Binding of a 20-mer peptide (20 uM) corresponding to the C-terminus of CD95 (Fas) to TIP-1 can be inhibited by an 8-mer peptide corresponding to the C-terminus of TAX. 50% inhibition can be achieved by 20-100 uM of inhibitor.

FIGURE 7 Binding of a 20-mer peptide (20 uM) corresponding to the C-terminus of TAX to TIP-1 can be inhibited by an 8-mer peptide corresponding to the C-terminus of CD95 (Fas). 50% inhibition can be achieved by 500 uM of inhibitor.

FIGURE 8 Binding of a 20-mer peptide (1 uM) corresponding to the C-terminus of BLR-1 (CXCR5) to KIAA0807 (PDZ domain)-GST fusion protein can be inhibited by an 8-mer peptide corresponding to the C-terminus of BLR-1 and a small molecule inhibitor (acetyl-LTTF). 50% inhibition can be achieved by greater than 100 uM of the 8-mer peptide and 1 uM of the small molecule inhibitor.

FIGURE 9 Binding of a 20-mer peptide (10 uM) corresponding to the C-

terminus of DOCK2 to KIAA0807 (PDZ domain)-GST fusion protein can be inhibited by an 8-mer peptide corresponding to the C-terminus of DOCK2 and a small molecule inhibitor (acetyl-STDL). 50% inhibition can be achieved by 250 uM of the 8-mer peptide and less than 250 uM of the small molecule inhibitor.

5 Tables

	Table 1	Amino Acid Classification
	Table 2	Protein-Ligand Pairs
	Table 3	PDZ Domains
	Table 3A	Note on Table 3
10	Table 4	PL Peptides
	Table 5	Exemplary PL Motifs
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	Table 7	PDZ Domain-Containing Genes Expressed in T Cells and B Cells
	Table 8	Linker sequences introduced in cloning
15	Table 9	Correlation of HPV E6 PL Motif and Oncogenic Activity

5. DEFINITIONS

5.1 A “fusion protein” or “fusion polypeptide” as used herein refers to a composite protein, i.e., a single contiguous amino acid sequence, made up of two (or more) distinct, heterologous polypeptides which are not normally fused together in a single amino acid sequence. Thus, a fusion protein can include a single amino acid sequence that contains two entirely distinct amino acid sequences or two similar or identical polypeptide sequences, provided that these sequences are not normally found together in the same configuration in a single amino acid sequence found in nature. Fusion proteins can generally be prepared using either recombinant nucleic acid methods, i.e., as a result of transcription and translation of a recombinant gene fusion product, which fusion comprises a segment encoding a polypeptide of the invention and a segment encoding a heterologous protein, or by chemical synthesis methods well known in the art.

5.2 A “fusion protein construct” as used herein is a polynucleotide encoding a fusion protein.

5.3 As used herein, the term “PDZ domain” refers to protein sequence (i.e., modular protein domain) of approximately 90 amino acids, characterized by homology to the brain synaptic protein PSD-95, the Drosophila septate junction protein Discs-Large (DLG), and the epithelial tight junction protein ZO1 (ZO1). PDZ domains are also known as Discs-Large

homology repeats (“DHRs”) and GLGF (SEQ ID NO: 402) repeats. PDZ domains generally appear to maintain a core consensus sequence (Doyle, D. A., 1996, *Cell* 85: 1067-76).

PDZ domains are found in diverse membrane-associated proteins including members of the MAGUK family of guanylate kinase homologs, several protein phosphatases and kinases, neuronal nitric oxide synthase, and several dystrophin-associated proteins, collectively known as syntrophins.

Exemplary PDZ domain-containing proteins and PDZ domain sequences are shown in TABLE 3. The term “PDZ domain” also encompasses variants (e.g., naturally occurring variants) of the sequences of TABLE 3 (e.g., polymorphic variants, variants with conservative substitutions, and the like). Typically, PDZ domains are substantially identical to those shown in TABLE 3, e.g., at least about 70%, at least about 80%, or at least about 90% amino acid residue identity when compared and aligned for maximum correspondence.

5.4 As used herein, the term “PDZ protein” refers to a naturally occurring protein containing a PDZ domain, e.g., a human protein. Exemplary PDZ proteins include CASK, MPP1, DLG1, PSD95, NeDLG, TAX33, SYN1a, TAX43, LDP, LIM, LIMK1, LIMK2, MPP2, NOS1, AF6, PTN-4, pIL16, 41.8kD, KIAA0559, RGS12, KIAA0316, DVL1, TAX40, TIAM1, MINT1, KIAA0303, CBP, MINT3, TAX2, KIAA0561. Exemplary PDZ proteins are listed in TABLE 2 and TABLE 3.

5.5 As used herein, the term “PDZ-domain polypeptide” refers to a polypeptide containing a PDZ domain, such as a fusion protein including a PDZ domain sequence, a naturally occurring PDZ protein, or an isolated PDZ domain peptide.

5.6 As used herein, the term “PL protein” or “PDZ Ligand protein” refers to a naturally occurring protein that forms a molecular complex with a PDZ-domain, or to a protein whose carboxy-terminus, when expressed separately from the full length protein (e.g., as a peptide fragment of 4-25 residues, e.g., 16 residues), forms such a molecular complex. The molecular complex can be observed *in vitro* using the “A assay” or “G assay” described *infra*, or *in vivo*. Exemplary PL proteins listed in TABLE 2 are demonstrated to bind specific PDZ proteins. This definition is not intended to include anti-PDZ antibodies and the like.

5.7 As used herein, a "PL sequence" refers to the amino acid sequence of the C-terminus of a PL protein (e.g., the C-terminal 2, 3, 4, 5, 6, 7, 8, 9, 10, 12, 14, 16, 20 or 25 residues) ("C-terminal PL sequence") or to an internal sequence known to bind a PDZ domain ("internal PL sequence").

5

5.8 As used herein, a "PL peptide" is a peptide of having a sequence from, or based on, the sequence of the C-terminus of a PL protein. Exemplary PL peptides (biotinylated) are listed in **TABLE 4**.

10 5.9 As used herein, a "PL fusion protein" is a fusion protein that has a PL sequence as one domain, typically as the C-terminal domain of the fusion protein. An exemplary PL fusion protein is a tat-PL sequence fusion.

15 5.10 As used herein, the term "PL inhibitor peptide sequence" refers to PL peptide amino acid sequence that (in the form of a peptide or PL fusion protein) inhibits the interaction between a PDZ domain polypeptide and a PL peptide (e.g., in an A assay or a G assay).

20 5.11 As used herein, a "PDZ-domain encoding sequence" means a segment of a polynucleotide encoding a PDZ domain. In various embodiments, the polynucleotide is DNA, RNA, single stranded or double stranded.

25 5.12 As used herein, the terms "antagonist" and "inhibitor," when used in the context of modulating a binding interaction (such as the binding of a PDZ domain sequence to a PL sequence), are used interchangeably and refer to an agent that reduces the binding of the, e.g., PL sequence (e.g., PL peptide) and the, e.g., PDZ domain sequence (e.g., PDZ protein, PDZ domain peptide).

30 5.13 As used herein, the terms "agonist" and "enhancer," when used in the context of modulating a binding interaction (such as the binding of a PDZ domain sequence to a PL sequence), are used interchangeably and refer to an agent that increases the binding of the, e.g., PL sequence (e.g., PL peptide) and the, e.g., PDZ domain sequence (e.g., PDZ protein, PDZ domain peptide).

5.14 As used herein, the terms "peptide mimetic," "peptidomimetic," and "peptide analog" are used interchangeably and refer to a synthetic chemical compound which has substantially the same structural and/or functional characteristics of an PL inhibitory or PL binding peptide of the invention. The mimetic can be either entirely composed of synthetic, non-natural analogues of amino acids, or, is a chimeric molecule of partly natural peptide amino acids and partly non-natural analogs of amino acids. The mimetic can also incorporate any amount of natural amino acid conservative substitutions as long as such substitutions also do not substantially alter the mimetic's structure and/or inhibitory or binding activity. As with polypeptides of the invention which are conservative variants, routine experimentation will determine whether a mimetic is within the scope of the invention, i.e., that its structure and/or function is not substantially altered. Thus, a mimetic composition is within the scope of the invention if it is capable of binding to a PDZ domain and/or inhibiting a PL-PDZ interaction.

Polypeptide mimetic compositions can contain any combination of nonnatural structural components, which are typically from three structural groups: a) residue linkage groups other than the natural amide bond ("peptide bond") linkages; b) non-natural residues in place of naturally occurring amino acid residues; or c) residues which induce secondary structural mimicry, i.e., to induce or stabilize a secondary structure, e.g., a beta turn, gamma turn, beta sheet, alpha helix conformation, and the like.

A polypeptide can be characterized as a mimetic when all or some of its residues are joined by chemical means other than natural peptide bonds. Individual peptidomimetic residues can be joined by peptide bonds, other chemical bonds or coupling means, such as, e.g., glutaraldehyde, N-hydroxysuccinimide esters, bifunctional maleimides, N,N'-dicyclohexylcarbodiimide (DCC) or N,N'-diisopropylcarbodiimide (DIC). Linking groups that can be an alternative to the traditional amide bond ("peptide bond") linkages include, e.g., ketomethylene (e.g., $-C(=O)-CH_2-$ for $-C(=O)-NH-$), aminomethylene (CH_2-NH), ethylene, olefin ($CH=CH$), ether (CH_2-O), thioether (CH_2-S), tetrazole (CN_4), thiazole, retroamide, thioamide, or ester (see, e.g., Spatola (1983) in *Chemistry and Biochemistry of Amino Acids, Peptides and Proteins*, Vol. 7, pp 267-357, A Peptide Backbone Modifications, Marcell Dekker, NY).

A polypeptide can also be characterized as a mimetic by containing all or some non-natural residues in place of naturally occurring amino acid residues. Nonnatural residues

are well described in the scientific and patent literature; a few exemplary nonnatural compositions useful as mimetics of natural amino acid residues and guidelines are described below.

Mimetics of aromatic amino acids can be generated by replacing by, e.g., D- or L- naphylalanine; D- or L- phenylglycine; D- or L-2 thieneylalanine; D- or L-1, -2, 3-, or 4-pyreneylalanine; D- or L-3 thieneylalanine; D- or L-(2-pyridinyl)-alanine; D- or L-(3-pyridinyl)-alanine; D- or L-(2-pyrazinyl)-alanine; D- or L-(4-isopropyl)-phenylglycine; D-(trifluoromethyl)-phenylglycine; D-(trifluoromethyl)-phenylalanine; D-p-fluorophenylalanine; D- or L-p-biphenylphenylalanine; K- or L-p-methoxybiphenylphenylalanine; D- or L-2-indole(alkyl)alanines; and, D- or L-alkylainines, where alkyl can be substituted or unsubstituted methyl, ethyl, propyl, hexyl, butyl, pentyl, isopropyl, iso-butyl, sec-isotyl, iso-pentyl, or a non-acidic amino acids. Aromatic rings of a nonnatural amino acid include, e.g., thiazolyl, thiophenyl, pyrazolyl, benzimidazolyl, naphthyl, furanyl, pyrrolyl, and pyridyl aromatic rings.

Mimetics of acidic amino acids can be generated by substitution by, e.g., non-carboxylate amino acids while maintaining a negative charge; (phosphono)alanine; sulfated threonine. Carboxyl side groups (e.g., aspartyl or glutamyl) can also be selectively modified by reaction with carbodiimides ($R=N-C-N=R$) such as, e.g., 1-cyclohexyl-3(2-morpholinyl-(4-ethyl) carbodiimide or 1-ethyl-3(4-azonia- 4,4- dimetholpentyl) carbodiimide. Aspartyl or glutamyl can also be converted to asparaginy and glutaminy residues by reaction with ammonium ions.

Mimetics of basic amino acids can be generated by substitution with, e.g., (in addition to lysine and arginine) the amino acids ornithine, citrulline, or (guanidino)-acetic acid, or (guanidino)alkyl-acetic acid, where alkyl is defined above. Nitrile derivative (e.g., containing the CN-moiety in place of COOH) can be substituted for asparagine or glutamine. Asparaginy and glutaminy residues can be deaminated to the corresponding aspartyl or glutamyl residues.

Arginine residue mimetics can be generated by reacting arginyl with, e.g., one or more conventional reagents, including, e.g., phenylglyoxal, 2,3-butanedione, 1,2-cyclohexanedione, or ninhydrin, preferably under alkaline conditions.

Tyrosine residue mimetics can be generated by reacting tyrosyl with, e.g., aromatic diazonium compounds or tetranitromethane. N-acetylimidizol and tetranitromethane can be used to form O-acetyl tyrosyl species and 3-nitro derivatives, respectively.

Cysteine residue mimetics can be generated by reacting cysteinyl residues with, e.g., alpha-haloacetates such as 2-chloroacetic acid or chloroacetamide and corresponding amines; to give carboxymethyl or carboxyamidomethyl derivatives. Cysteine residue mimetics can also be generated by reacting cysteinyl residues with, e.g., bromo-trifluoroacetone, alpha-bromo-beta-(5-imidozoyl) propionic acid; chloroacetyl phosphate, N-alkylmaleimides, 3-nitro-2-pyridyl disulfide; methyl 2-pyridyl disulfide; p-chloromercuribenzoate; 2-chloromercuri-4-nitrophenol; or, chloro-7-nitrobenzo-oxa-1,3-diazole.

Lysine mimetics can be generated (and amino terminal residues can be altered) by reacting lysinyl with, e.g., succinic or other carboxylic acid anhydrides. Lysine and other alpha-amino-containing residue mimetics can also be generated by reaction with imidoesters, such as methyl picolinimide, pyridoxal phosphate, pyridoxal, chloroborohydride, trinitrobenzenesulfonic acid, O-methylisourea, 2,4, pentanedione, and transamidase-catalyzed reactions with glyoxylate.

Mimetics of methionine can be generated by reaction with, e.g., methionine sulfoxide. Mimetics of proline include, e.g., pipecolic acid, thiazolidine carboxylic acid, 3- or 4- hydroxy proline, dehydropoline, 3- or 4-methylproline, or 3,3,-dimethylproline. Histidine residue mimetics can be generated by reacting histidyl with, e.g., diethylprocarbonate or para-bromophenacyl bromide.

Other mimetics include, e.g., those generated by hydroxylation of proline and lysine; phosphorylation of the hydroxyl groups of seryl or threonyl residues; methylation of the alpha-amino groups of lysine, arginine and histidine; acetylation of the N-terminal amine; methylation of main chain amide residues or substitution with N-methyl amino acids; or amidation of C-terminal carboxyl groups.

A component of a natural polypeptide (e.g., a PL polypeptide or PDZ polypeptide) can also be replaced by an amino acid (or peptidomimetic residue) of the opposite chirality. Thus, any amino acid naturally occurring in the L-configuration (which can also be referred to as the R or S, depending upon the structure of the chemical entity) can be replaced with the amino acid of the same chemical structural type or a peptidomimetic, but of the opposite chirality, generally referred to as the D- amino acid, but which can additionally be referred to as the R- or S- form.

The mimetics of the invention can also include compositions that contain a structural mimetic residue, particularly a residue that induces or mimics secondary structures,



such as a beta turn, beta sheet, alpha helix structures, gamma turns, and the like. For example, substitution of natural amino acid residues with D-amino acids; N-alpha-methyl amino acids; C-alpha-methyl amino acids; or dehydroamino acids within a peptide can induce or stabilize beta turns, gamma turns, beta sheets or alpha helix conformations. Beta turn mimetic structures have been described, e.g., by Nagai (1985) Tet. Lett. 26:647-650; Feigl (1986) J. Amer. Chem. Soc. 108:181-182; Kahn (1988) J. Amer. Chem. Soc. 110:1638-1639; Kemp (1988) Tet. Lett. 29:5057-5060; Kahn (1988) J. Molec. Recognition 1:75-79. Beta sheet mimetic structures have been described, e.g., by Smith (1992) J. Amer. Chem. Soc. 114:10672-10674. For example, a type VI beta turn induced by a cis amide surrogate, 1,5-disubstituted tetrazol, is described by Beusen (1995) Biopolymers 36:181-200. Incorporation of achiral omega-amino acid residues to generate polymethylene units as a substitution for amide bonds is described by Banerjee (1996) Biopolymers 39:769-777. Secondary structures of polypeptides can be analyzed by, e.g., high-field 1H NMR or 2D NMR spectroscopy, see, e.g., Higgins (1997) J. Pept. Res. 50:421-435. See also, Hruby (1997) Biopolymers 43:219-266, Balaji, et al., U.S. Pat. No. 5,612,895.

5.15 As used herein, "peptide variants" and "conservative amino acid substitutions" refer to peptides that differ from a reference peptide (e.g., a peptide having the sequence of the carboxy-terminus of a specified PL protein) by substitution of an amino acid residue having similar properties (based on size, polarity, hydrophobicity, and the like). Thus, insofar as the compounds that are encompassed within the scope of the invention are partially defined in terms of amino acid residues of designated classes, the amino acids may be generally categorized into three main classes: hydrophilic amino acids, hydrophobic amino acids and cysteine-like amino acids, depending primarily on the characteristics of the amino acid side chain. These main classes may be further divided into subclasses. Hydrophilic amino acids include amino acids having acidic, basic or polar side chains and hydrophobic amino acids include amino acids having aromatic or apolar side chains. Apolar amino acids may be further subdivided to include, among others, aliphatic amino acids. The definitions of the classes of amino acids as used herein are as follows:

"Hydrophobic Amino Acid" refers to an amino acid having a side chain that is uncharged at physiological pH and that is repelled by aqueous solution. Examples of genetically encoded hydrophobic amino acids include Ile, Leu and Val. Examples of non-

genetically encoded hydrophobic amino acids include t-BuA.

"Aromatic Amino Acid" refers to a hydrophobic amino acid having a side chain containing at least one ring having a conjugated π -electron system (aromatic group). The aromatic group may be further substituted with groups such as alkyl, alkenyl, alkynyl, hydroxyl, sulfanyl, nitro and amino groups, as well as others. Examples of genetically encoded aromatic amino acids include Phe, Tyr and Trp. Commonly encountered non-genetically encoded aromatic amino acids include phenylglycine, 2-naphthylalanine, β -2-thienylalanine, 1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid, 4-chloro-phenylalanine, 2-fluorophenyl-alanine, 3-fluorophenylalanine and 4-fluorophenylalanine.

"Apolar Amino Acid" refers to a hydrophobic amino acid having a side chain that is generally uncharged at physiological pH and that is not polar. Examples of genetically encoded apolar amino acids include Gly, Pro and Met. Examples of non-encoded apolar amino acids include Cha.

"Aliphatic Amino Acid" refers to an apolar amino acid having a saturated or unsaturated straight chain, branched or cyclic hydrocarbon side chain. Examples of genetically encoded aliphatic amino acids include Ala, Leu, Val and Ile. Examples of non-encoded aliphatic amino acids include Nle.

"Hydrophilic Amino Acid" refers to an amino acid having a side chain that is attracted by aqueous solution. Examples of genetically encoded hydrophilic amino acids include Ser and Lys. Examples of non-encoded hydrophilic amino acids include Cit and hCys.

"Acidic Amino Acid" refers to a hydrophilic amino acid having a side chain pK value of less than 7. Acidic amino acids typically have negatively charged side chains at physiological pH due to loss of a hydrogen ion. Examples of genetically encoded acidic amino acids include Asp and Glu.

"Basic Amino Acid" refers to a hydrophilic amino acid having a side chain pK value of greater than 7. Basic amino acids typically have positively charged side chains at physiological pH due to association with hydronium ion. Examples of genetically encoded basic amino acids include Arg, Lys and His. Examples of non-genetically encoded basic amino acids include the non-cyclic amino acids ornithine, 2,3-diaminopropionic acid, 2,4-diaminobutyric acid and homoarginine.

"Polar Amino Acid" refers to a hydrophilic amino acid having a side chain that is uncharged at physiological pH, but which has a bond in which the pair of electrons shared

in common by two atoms is held more closely by one of the atoms. Examples of genetically encoded polar amino acids include Asx and Glx. Examples of non-genetically encoded polar amino acids include citrulline, N-acetyl lysine and methionine sulfoxide.

"Cysteine-Like Amino Acid" refers to an amino acid having a side chain capable of forming a covalent linkage with a side chain of another amino acid residue, such as a disulfide linkage. Typically, cysteine-like amino acids generally have a side chain containing at least one thiol (SH) group. Examples of genetically encoded cysteine-like amino acids include Cys. Examples of non-genetically encoded cysteine-like amino acids include homocysteine and penicillamine.

As will be appreciated by those having skill in the art, the above classification are not absolute -- several amino acids exhibit more than one characteristic property, and can therefore be included in more than one category. For example, tyrosine has both an aromatic ring and a polar hydroxyl group. Thus, tyrosine has dual properties and can be included in both the aromatic and polar categories. Similarly, in addition to being able to form disulfide linkages, cysteine also has apolar character. Thus, while not strictly classified as a hydrophobic or apolar amino acid, in many instances cysteine can be used to confer hydrophobicity to a peptide.

Certain commonly encountered amino acids which are not genetically encoded of which the peptides and peptide analogues of the invention may be composed include, but are not limited to, β -alanine (b-Ala) and other omega-amino acids such as 3-aminopropionic acid (Dap), 2,3-diaminopropionic acid (Dpr), 4-aminobutyric acid and so forth; α -aminoisobutyric acid (Aib); ϵ -aminohexanoic acid (Aha); δ -aminovaleric acid (Ava); N-methylglycine or sarcosine (MeGly); ornithine (Orn); citrulline (Cit); t-butylalanine (t-BuA); t-butylglycine (t-BuG); N-methylisoleucine (MeIle); phenylglycine (Phg); cyclohexylalanine (Cha); norleucine (Nle); 2-naphthylalanine (2-Nal); 4-chlorophenylalanine (Phe(4-Cl)); 2-fluorophenylalanine (Phe(2-F)); 3-fluorophenylalanine (Phe(3-F)); 4-fluorophenylalanine (Phe(4-F)); penicillamine (Pen); 1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid (Tic); β -2-thienylalanine (Thi); methionine sulfoxide (MSO); homoarginine (hArg); N-acetyl lysine (AcLys); 2,3-diaminobutyric acid (Dab); 2,3-diaminobutyric acid (Dbu); p-aminophenylalanine (Phe(pNH₂)); N-methyl valine (MeVal); homocysteine (hCys) and homoserine (hSer). These amino acids also fall conveniently into the categories defined above.

The classifications of the above-described genetically encoded and non-encoded amino acids are summarized in **TABLE 1**, below. It is to be understood that **TABLE 1** is for illustrative purposes only and does not purport to be an exhaustive list of amino acid residues which may comprise the peptides and peptide analogues described herein. Other amino acid residues which are useful for making the peptides and peptide analogues described herein can be found, e.g., in Fasman, 1989, CRC Practical Handbook of Biochemistry and Molecular Biology, CRC Press, Inc., and the references cited therein. Amino acids not specifically mentioned herein can be conveniently classified into the above-described categories on the basis of known behavior and/or their characteristic chemical and/or physical properties as compared with amino acids specifically identified.

TABLE 1

Classification	Genetically Encoded	Genetically Non-Encoded
Hydrophobic		
Aromatic	F, Y, W	Phg, Nal, Thi, Tic, Phe(4-Cl), Phe(2-F), Phe(3-F), Phe(4-F), Pyridyl Ala, Benzothienyl Ala
Apolar	M, G, P	
Aliphatic	A, V, L, I	t-BuA, t-BuG, Melle, Nle, MeVal, Cha, bAla, MeGly, Aib
Hydrophilic		
Acidic	D, E	
Basic	H, K, R	Dpr, Orn, hArg, Phe(p-NH ₂), DBU, A ₂ BU
Polar	Q, N, S, T, Y	Cit, AcLys, MSO, hSer
Cysteine-Like	C	Pen, hCys, p-methyl Cys

5.16 As used herein, a “detectable label” has the ordinary meaning in the art and refers to an atom (e.g., radionuclide), molecule (e.g., fluorescein), or complex, that is or can be used to detect (e.g., due to a physical or chemical property), indicate the presence of a molecule or to enable binding of another molecule to which it is covalently bound or otherwise associated. The term “label” also refers to covalently bound or otherwise associated molecules (e.g., a biomolecule such as an enzyme) that act on a substrate to produce a detectable atom, molecule or complex. Detectable labels suitable for use in the present invention include any composition detectable by spectroscopic, photochemical, biochemical, immunochemical,



electrical, optical or chemical means. Labels useful in the present invention include biotin for staining with labeled streptavidin conjugate, magnetic beads (e.g., Dynabeads™), fluorescent dyes (e.g., fluorescein, Texas red, rhodamine, green fluorescent protein, enhanced green fluorescent protein, and the like), radiolabels (e.g., ^3H , ^{125}I , ^{35}S , ^{14}C , or ^{32}P), enzymes (e.g., hydrolases, particularly phosphatases such as alkaline phosphatase, esterases and glycosidases, or oxidoreductases, particularly peroxidases such as horse radish peroxidase, and others commonly used in ELISAs), substrates, cofactors, inhibitors, chemiluminescent groups, chromogenic agents, and colorimetric labels such as colloidal gold or colored glass or plastic (e.g., polystyrene, polypropylene, latex, etc.) beads. Patents teaching the use of such labels include U.S. Patent Nos. 3,817,837; 3,850,752; 3,939,350; 3,996,345; 4,277,437; 4,275,149; and 4,366,241. Means of detecting such labels are well known to those of skill in the art. Thus, for example, radiolabels and chemiluminescent labels may be detected using photographic film or scintillation counters, fluorescent markers may be detected using a photodetector to detect emitted light (e.g., as in fluorescence-activated cell sorting). Enzymatic labels are typically detected by providing the enzyme with a substrate and detecting the reaction product produced by the action of the enzyme on the substrate, and colorimetric labels are detected by simply visualizing the colored label. Thus, a label is any composition detectable by spectroscopic, photochemical, biochemical, immunochemical, electrical, optical or chemical means. The label may be coupled directly or indirectly to the desired component of the assay according to methods well known in the art. Non-radioactive labels are often attached by indirect means. Generally, a ligand molecule (e.g., biotin) is covalently bound to the molecule. The ligand then binds to an anti-ligand (e.g., streptavidin) molecule which is either inherently detectable or covalently bound to a signal generating system, such as a detectable enzyme, a fluorescent compound, or a chemiluminescent compound. A number of ligands and anti-ligands can be used. Where a ligand has a natural anti-ligand, for example, biotin, thyroxine, and cortisol, it can be used in conjunction with the labeled, naturally occurring anti-ligands. Alternatively, any haptenic or antigenic compound can be used in combination with an antibody. The molecules can also be conjugated directly to signal generating compounds, e.g., by conjugation with an enzyme or fluorophore. Means of detecting labels are well known to those of skill in the art. Thus, for example, where the label is a radioactive label, means for detection include a scintillation counter, photographic film as in autoradiography, or storage phosphor imaging. Where the label is a fluorescent label, it may be detected by exciting the fluorochrome with the

appropriate wavelength of light and detecting the resulting fluorescence. The fluorescence may be detected visually, by means of photographic film, by the use of electronic detectors such as charge coupled devices (CCDs) or photomultipliers and the like. Similarly, enzymatic labels may be detected by providing the appropriate substrates for the enzyme and detecting the resulting reaction product. Also, simple colorimetric labels may be detected by observing the color associated with the label. It will be appreciated that when pairs of fluorophores are used in an assay, it is often preferred that they have distinct emission patterns (wavelengths) so that they can be easily distinguished.

5.17 As used herein, the term "substantially identical" in the context of comparing amino acid sequences, means that the sequences have at least about 70%, at least about 80%, or at least about 90% amino acid residue identity when compared and aligned for maximum correspondence. An algorithm that is suitable for determining percent sequence identity and sequence similarity is the FASTA algorithm, which is described in Pearson, W.R. & Lipman, D.J., 1988, *Proc. Natl. Acad. Sci. U.S.A.* 85: 2444. See also *W. R. Pearson, 1996, Methods Enzymol.* 266: 227-258. Preferred parameters used in a FASTA alignment of DNA sequences to calculate percent identity are optimized, BL50 Matrix 15: -5, k-tuple = 2; joining penalty = 40, optimization = 28; gap penalty -12, gap length penalty = -2; and width = 16.

5.18 As used herein, "hematopoietic cells" include leukocytes including lymphocytes (T cells, B cells and NK cells), monocytes, and granulocytes (i.e., neutrophils, basophils and eosinophils), macrophages, dendritic cells, megakaryocytes, reticulocytes, erythrocytes, and CD34⁺ stem cells.

5.19 As used herein, the terms "test compound" or "test agent" are used interchangeably and refer to a candidate agent that may have enhancer/agonist, or inhibitor/antagonist activity, e.g., inhibiting or enhancing an interaction such as PDZ-PL binding. The candidate agents or test compounds may be any of a large variety of compounds, both naturally occurring and synthetic, organic and inorganic, and including polymers (e.g., oligopeptides, polypeptides, oligonucleotides, and polynucleotides), small molecules, antibodies (as broadly defined herein), sugars, fatty acids, nucleotides and nucleotide analogs, analogs of naturally occurring structures (e.g., peptide mimetics, nucleic acid analogs, and the

like), and numerous other compounds. In certain embodiment, test agents are prepared from diversity libraries, such as random or combinatorial peptide or non-peptide libraries. Many libraries are known in the art that can be used, e.g., chemically synthesized libraries, recombinant (e.g., phage display libraries), and *in vitro* translation-based libraries. Examples of chemically synthesized libraries are described in Fodor et al., 1991, *Science* 251:767-773; Houghten et al., 1991, *Nature* 354:84-86; Lam et al., 1991, *Nature* 354:82-84; Medynski, 1994, *Bio/Technology* 12:709-710; Gallop et al., 1994, *J. Medicinal Chemistry* 37(9):1233-1251; Ohlmeyer et al., 1993, *Proc. Natl. Acad. Sci. USA* 90:10922-10926; Erb et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:11422-11426; Houghten et al., 1992, *Biotechniques* 13:412; Jayawickreme et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:1614-1618; Salmon et al., 1993, *Proc. Natl. Acad. Sci. USA* 90:11708-11712; PCT Publication No. WO 93/20242; and Brenner and Lerner, 1992, *Proc. Natl. Acad. Sci. USA* 89:5381-5383. Examples of phage display libraries are described in Scott and Smith, 1990, *Science* 249:386-390; Devlin et al., 1990, *Science*, 249:404-406; Christian, R.B., et al., 1992, *J. Mol. Biol.* 227:711-718; Lenstra, 1992, *J. Immunol. Meth.* 152:149-157; Kay et al., 1993, *Gene* 128:59-65; and PCT Publication No. WO 94/18318 dated August 18, 1994. *In vitro* translation-based libraries include but are not limited to those described in PCT Publication No. WO 91/05058 dated April 18, 1991; and Mattheakis et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:9022-9026. By way of examples of nonpeptide libraries, a benzodiazepine library (see e.g., Bunin et al., 1994, *Proc. Natl. Acad. Sci. USA* 91:4708-4712) can be adapted for use. Peptoid libraries (Simon et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:9367-9371) can also be used. Another example of a library that can be used, in which the amide functionalities in peptides have been permethylated to generate a chemically transformed combinatorial library, is described by Ostresh et al. (1994, *Proc. Natl. Acad. Sci. USA* 91:11138-11142).

5.20 The term "specific binding" refers to binding between two molecules, for example, a ligand and a receptor, characterized by the ability of a molecule (ligand) to associate with another specific molecule (receptor) even in the presence of many other diverse molecules, i.e., to show preferential binding of one molecule for another in a heterogeneous mixture of molecules. Specific binding of a ligand to a receptor is also evidenced by reduced binding of a detectably labeled ligand to the receptor in the presence of excess unlabeled ligand (i.e., a binding competition assay).



5.21 As used herein, a “plurality” of PDZ proteins (or corresponding PDZ domains or PDZ fusion polypeptides) has its usual meaning. In some embodiments, the plurality is at least 5, and often at least 25, at least 40, or at least 60 different PDZ proteins. In some embodiments, the plurality is selected from the list of PDZ polypeptides listed in **Table 2** or **Table 7**. In some embodiments, the plurality of different PDZ proteins are from (i.e., expressed in) a particular specified tissue or a particular class or type of cell. In some embodiments, the plurality of different PDZ proteins represents a substantial fraction (e.g., typically at least 50%, more often at least 80%) of all of the PDZ proteins known to be, or suspected of being, expressed in the tissue or cell(s), e.g., all of the PDZ proteins known to be present in lymphocytes or hematopoietic cells. In some embodiments, the plurality is at least 50%, usually at least 80%, at least 90% or all of the PDZ proteins disclosed herein as being expressed in hematopoietic cells (see Tables 2 and 6). In an embodiment, the plurality includes at least 1, often at least 2, sometimes at least 5 or at least 10 and sometimes all of the following PDZ proteins: BAI I associated protein, Connector enhancer, DLG5 (pdlg), DVL3, GTPase, Guanine-exchange factor 1, PDZ domain containing prot., KIAA147, KIAA0300, KIAA0380, KIAA0440, KIAA0545, KIAA0807, KIAA0858, KIAA0902, novel serine protease, PDZK1, PICK8, PTN-3, RPIP8, serine protease, 26s subunit p27, hSYNTENIN, TAX1-IP, TAX2-like protein, wwp3, X11 prot. beta, ZO1. When referring to PL ligands or corresponding PL proteins (e.g., corresponding to those listed in Table 2, Table 4, Table 5, or elsewhere herein) a “plurality” may refer to at least 5, at least 10, and often at least 25 PLs such as those specifically listed herein, or to the classes and percentages set forth *supra* for PDZ domains.

6. DETAILED DESCRIPTION OF THE INVENTION

The present inventors have discovered that interactions between PDZ proteins and PL proteins play an important and extensive role in the biological function of hematopoietic cells and other cells involved in the immune response. Although PDZ-PL interactions were known in the nervous system (i.e., in neurons), their universal importance in hematopoietic cell function, especially in function of T cells and B cells, and their fundamental role in modulation of the immune response has not been recognized. In particular, the present



inventors have surprisingly discovered that cell adhesion molecules that mediate cell-cell interaction in the hematopoietic system are PDZ-binding proteins (PL proteins) and bind to PDZ proteins. The inventors have identified numerous interactions between PDZ proteins and PL proteins present in immune system cells, and the invention provides reagents and methods for affecting biological function in the immune system by inhibiting these interactions. As used herein, the term “biological function” in the context of a cell, refers to a detectable biological activity normally carried out by the cell, e.g., a phenotypic change such as proliferation, cell activation (e.g., T cell activation, B cell activation, T-B cell conjugate formation), cytokine release, degranulation, tyrosine phosphorylation, ion (e.g., calcium) flux, metabolic activity, apoptosis, changes in gene expression, maintenance of cell structure, cell migration, adherence to a substrate, signal transduction, cell-cell interactions, and others described herein or known in the art.

In one aspect, the present invention relates to peptides, peptide analogues or mimetics, pharmaceutical compositions, and methods of using such compositions to regulate the biological activities of hematopoietic cells, e.g. T cells and B cells, or other cells (e.g., endothelial cells) that necessary for immune function. The invention further relates to methods of using the compositions to modulate hematopoietic cell activation and immune function, as well as assays for such inhibitors.

TABLE 2 summarizes an extensive analysis of protein interactions in T cells and B cells. PDZ proteins, the vast majority of which were not previously known to be expressed in immune system cells, are listed in the top row of **TABLE 2**. The first column of the table lists PL proteins. Positions in the matrix denoted by the letter “A,” “G,” “G’,” or “G’” indicate that an interaction between the PDZ protein and the PL has been detected in novel binding assays (described in detail *infra*). A blank cell indicates that no interaction was detected using the assays of the invention. (S) – Indicates “sticky” peptide ligand having high background signal (i.e., in one or more versions of the G assay, the signal of ligand binding in the GST alone background wells was repeatedly above 0.5 OD units). An asterisk (*) denotes a PL-PDZ interaction previously reported in the scientific literature.

PDZ-GST fusion Protein:			CASK	MPP1	LIMK1	K303	K807	DLG1	PSD95
PDZ LIGAND	CODE	SEQ							
CD6	AA6L	ISAA							
CD49E (alpha-4)	AA11L	TSDA							
CD49F (Aform, alpha6)	AA12L	TSDA							
CD105 (endoglin)	AA16L	SSMA							
CD166 (CD6L)	AA20L	KTEA							
CC CKR-2	AA42L	KEGA							
CD138 (syndecan-1)	AA18L	EFYA	*						
Syndecan-2 (S)	AA39L	EFYA							
CD148 (DEP-1)	AA19L	GYIA							
CD98 (2F4) (S)	AA15L	PYAA							
CLASP-1	AA1L	SAEV						G	A
CLASP-4	AA3L-V	YAEV						A	A
NMDA	AA34.2L	ESDV		A	A			A/G	A/G
VCAM1	AA17L	KSKV		A		A	G'/G*	A	
CLASP-2	AA2L	SSVV						A/G	A/G
CD95 (Apo-1/Fas)	AA13L	QSLV						A/G/G'	A/G/G'
Spectrin beta (S)	AA32L	VSVF		G*		G*	G'/G*	G'/G*	G'/G*
KV1.3	AA33L	FTDV			A		G'/G*	*A/G/G'/G*	*A/G/G'/G*
DNAM-1	AA22L	KTRV		A				A	A/G/G'
Neurexin 3	AA36L	TTRV					G*		
TAX	AA56L	ETEV					G'	G'/G*	G'/G*
CD83	AA47L	TELV						A	A
CD44 (long form)	AA9L	KIGV		G					
Neurexin (S)	AA38L	EYV	G*	A*	A			A/G	A/G
CD97 (CD55L)	AA14L	ESGI						A	
CD38 (S)	AA8L	TSEI					G'		G'
Mannose receptor	AA31L	HSVI							
Glycophorin C	AA37L	EYFI		*				G	G
Galectin3	AA26L	YTMI							
CDw128A (IL8RA)	AA29.1L	SSNL						A	
CD3n	AA4L	SSQL					G*	A	A
LPAP	AA30L	VTAL					G'/G*	A	
CD46 (form 1)	AA10L	FTSL					G'/G*	A/G	A/G
CDw128B (IL8RB)	AA29.2L	STTL					G'/G*	A/G	A
DOCK2	AA40L	STD					G'/G*	A	A/G
PAG	AA58L	ITRL					G'		
CD34	AA7L	DTEL					G'/G*	A	A
CD5	AA49L	AQRL							
CC CKR-4	AA44L	HDAL							
FceRIb	AA25L	PIDL							
CDw137 (4-1BB-IL4) (S)	AA21L	GCEL							
FasLigand	AA23L-M	LYKL							
CD62E	AA48L	SYIL							
CC CKR-1R	AA41L	SAGF							
CDw125 (IL5R)	AA28L	DSVF							
BLR-1	AA45L	LTTF					G'		
CC CKR-3	AA43L	SIVF							
CD114 (G-CSFR)	AA27L	LGSF							
V-gated Ca2+ channel (S)	AA46L	DHWC							
PDZ-GST fusion Protein:			CASK	MPP1	LIMK1	K303	K807	DLG1	PSD95

Table 2

PDZ-GST fusion Protein:	NeDLG	SNTa1	TAX-IP43	LDP	LIM	MINT1	X11β	K440	K545	TAX-IP2
PDZ LIGAND										
CD6										
CD49E (alpha-4)										
CD49F (Aform, alpha6)										
CD105 (endoglin)					G'				G"	
CD166 (CD6L)										
CC CKR-2										
CD138 (syndecan)										
Syndecan-2 (S)										
CD148 (DEP-1)										
CD98 (2F4) (S)										
CLASP-1	G									
CLASP-4	A			A		A				
NMDA	A/G	G	A			A/G				A
VCAM1	A			A						
CLASP-2	A/G/G'									
CD95 (Apo-1/Fas)	A/G/G'									
Spectrin beta (S)	G'/G"		G'							
KV1.3	A/G/G'/G"					G/G'				
DNAM-1	A				A					
Neuroigin 3										
TAX	G'/G"		G'/G"							
CD83	A									
CD44 (long form)						G				
Neurexin (S)	G	A	A			A/G				
CD97 (CD55L)										
CD38 (S)	G'		G'	G'						
Mannose receptor										
Glycophorin C	G					A				
Galectin3										
CDw128A (IL8RA)	A									
CD3h						A/G/G'				
LPAP						G				
CD46 (form 1)	G									
CDw128B (IL8RB)	A/G/G'									
DOCK2	G	G								
PAG										
CD34	G									
CD5										
CC CKR-4										
FceR1b						A/G'				
CDw137 (4-1BB ILA) (S)										
FasLigand										
CD62E										
CC CKR-1R										
CDw125 (IL5R)										
BLR-1						G				
CC CKR-3										
CD114 (G-CSFR)										
V-gated Ca2+ channel (S)										
PDZ-GST fusion Protein:	NeDLG	SYN1a	TAX-IP43	LDP	LIM	MINT1	X11β	K440	K545	TAX-IP2

Table 2

DZ-GST fusion Protein:	TAX-IP2L	TAX-IP33	MPP2	MINT3	TIP1	PTN-4	prIL16	CBP	41.8	K559
DZ LIGAND										
CD6									A	
CD49E (alpha-4)									A/G	
CD49F (Aform, alpha6)									A/G	
CD105 (endoglin)										
CD166 (CD6L)										
CC CKR-2										
CD138 (syndecan)									A/G	
Syndecan-2 (S)										
CD148 (DEP-1)										
CD98 (2F4) (S)			G							
CLASP-1										
CLASP-4									A	
NMDA			G			A/G			A/G	
VCAM1									A	
CLASP-2									A	
CD95 (Apo-1/Fas)					G'				A/G	
Spectrin beta (S)						G'		G'		
KV1.3					G'				A	
DNAM-1			G						A	
Neuroigin 3										
TAX					G'	G'				
CD83										
CD44 (long form)							G			
Neurexin (S)							A		A	
CD97 (CD55L)									A	
CD38 (S)										
Mannose receptor										
Glycophorin C			A						A	
Galectin3										
CDw128A (IL8RA)										
CD3n					G'/G'				A/G	
LPAP					G'					
CD46 (form 1)										
CDw128B (IL8RB)									A	
DOCK2										
PAG										
CD34										
CD5										
CC CKR-4										
FceR1b										
CDw137 (4-1BB ILA) (S)										
FasLigand										
CD62E										
CC CKR-1R										
CDw125 (IL5R)						G				
BLR-1										
CC CKR-3										
CD114 (G-CSFR)										
V-gated Ca2+ channel (S)										
PDZ-GST fusion Protein:	TAX-IP2L	TAX-IP33	MPP2	MINT3	TIP1	PTN-4	prIL16	CBP	41.8	K559

Table 2

DZ-GST fusion Protein:	AF6	PICK1	RGS12	PDZK1	K316	DLG5	Synt	WWP3	TAX-IP40	K858
DZ LIGAND										
CD6										
CD49E (alpha-4)										
CD49F (Aform, alpha6)										
CD105 (endoglin)										
CD166 (CD6L)										
CC CKR-2										
CD138 (syndecan)										
Syndecan-2 (S)										
CD148 (DEP-1)										
CD98 (2F4) (S)										
CLASP-1										
CLASP-4	A									
MDA			AVG							
VCAM1								G'		
CLASP-2										
CD95 (Apo-1/Fas)										
Spectrin beta (S)	G'/G"					G'		G'/G"		
KV1.3			A							
DNAM-1	A		A					G'/G"		
Neuroigin 3								G"		
TAX						G'/G"		G'		
CD83										
CD44 (long form)							G'			
Neurexin (S)	A		A						A	
CD97 (CD55L)										
CD38 (S)										
Mannose receptor										
Glycophorin C	A									
Galectin3										
CDw128A (IL8RA)										
CD3n										
LPAP										
CD46 (form 1)										
CDw128B (IL8RB)			*							
DOCK2										
PAG										
CD34										
CD5										
CC CKR-4										
FceRIb										
CDw137 (4-1BB ILA) (S)										
FasLigand										
CD62E										
CC CKR-1R										
CDw125 (IL5R)			G							
BLR-1				G'/G"						
CC CKR-3										
CD114 (G-CSFR)										
V-gated Ca2+ channel (S)										
PDZ-GST fusion Protein:	AF6	PICK1	RGS12	PDZK1	K316	DLG5	Synt	WWP3	TAX-IP40	K858

Table 2

PDZ-GST fusion Protein:	TIAM1	SP short	ConEn	DVL1	NSP	GEF	K902	K561	NOS1	LIMK2
PDZ LIGAND										
CD6										
CD49E (alpha-4)										
CD49F (Aform, alpha6)										
CD105 (endoglin)										
CD166 (CD6L)										
CC CKR-2										
CD138 (syndecan)	A									
Syndecan-2 (S)										
CD148 (DEP-1)										
CD98 (2F4) (S)										
CLASP-1										
CLASP-4										
NMDA				A				G		
VCAM1	A									
CLASP-2										
CD95 (Apo-1/Fas)										
Spectrin beta (S)										
KV1.3				A						
DNAM-1										
Neuroigin 3										
TAX										
CD83										
CD44 (long form)										
Neurexin (S)	A			A						
CD97 (CD55L)										
CD38 (S)										
Mannose receptor										
Glycophorin C										
Galectin3										
CDw128A (IL8RA)										
CD3n										
LPAP										
CD46 (form 1)										
CDw128B (IL8RB)										
DOCK2								G		
PAG										
CD34										
CD5										
CC CKR-4										
FceRIb										
CDw137 (4-1BB ILA) (S)										
FasLigand								G		
CD62E										
CC CKR-1R										
CDw125 (IL5R)										
BLR-1								G		
CC CKR-3										
CD114 (G-CSFR)										
V-gated Ca2+ channel (S)										
PDZ-GST fusion Protein:	TIAM1	SP short	ConEn	DVL1	NSP	GEF	K902	K561	NOS1	LIMK2

Table 2

As discussed in detail herein, the PDZ proteins listed in **TABLE 2** are naturally occurring proteins containing a PDZ domain. The present invention is particularly directed to the detection and modulation of interactions between PDZ proteins and PL proteins in hematopoietic cells. Exemplary PL proteins are listed in **TABLE 2**. Notably, as discussed

5 *infra*, many of these PL proteins have not previously been recognized as such in any cell system. A variety of PL protein classes are known, and the PL proteins described herein can be characterized as (1) "PL adhesion proteins" (2) "PL ion channel proteins" (3) "PL adaptor proteins" (4) "PL intracellular proteins" and (5) "PL cytokine receptor proteins."

As used herein, an adhesion protein is a cell surface protein involved in cell-cell

10 interaction by direct contact with cell surface molecules (e.g., transmembrane proteins or surface proteins) on a different cell. Thus, when a cell expressing a PL adhesion protein contacts an appropriate other cell, the PL adhesion protein localizes at the interface of the two cells and directly contacts a cell surface molecule on the second cell. A cell-cell interface is a region where the plasma membranes of two different cells are in close (generally <10 nm,

15 often about 1 nm) apposition. Typically, direct molecular contact means interaction of molecules at distances where Van der Waals forces are significant, generally less than about 1 nm. Exemplary PL adhesion proteins include CD6; CD49E (alpha-4); CD49F (a form, alpha6); CD138 (syndecan); CLASP-1; CLASP-4; VCAM1; CLASP-2; DNAM-1; CD83; CD44 (long form); CD97; (CD55L); CD3 η ; DOCK2; CD34; and Fc ϵ R1b. Thus, in one embodiment, the

20 PL proteins of the invention are PL adhesion proteins. In an embodiment, the invention provides methods and reagents, as detailed herein, for inhibiting interactions between PL adhesion proteins and PDZ proteins to modulate an immune response. In an embodiment, the inhibition or modulation occurs in a hematopoietic cell. In a related embodiment, the inhibition or modulation occurs in an endothelial cell. In a related embodiment, the inhibition or

25 modulation occurs in an endothelial cell. In a related embodiment, the inhibition or modulation occurs in an epithelial cells, keratinocytes, hepatocytes, cardiac myocytes.

As used herein, an ion channel protein means a transmembrane protein that itself catalyzes the passage of an ion from aqueous solution on one side of a lipid bilayer membrane to aqueous solution on the other side (e.g., by forming a small pore in the membrane). One

30 exemplary PL ion channel proteins is Kv1.3. Thus, in one embodiment, the PL proteins of the invention are PL ion channel proteins. In an embodiment, the invention provides methods and reagents, as detailed herein, for inhibiting interactions between PL ion channel proteins and

PDZ proteins to modulate an immune response. In an embodiment, the inhibition or modulation occurs in a hematopoietic cell. In a related embodiment, the inhibition or modulation occurs in an endothelial cell.

As used herein, an intercellular (i.e., cytosolic) protein has the normal meaning in the art and refers to a protein that is not membrane bound, e.g., has no transmembrane domain. Thus, in one embodiment, the PL proteins of the invention are PL intercellular proteins. Exemplary PL intercellular proteins include Glycophorin C and LPAP. In an embodiment, the invention provides methods and reagents, as detailed herein, for inhibiting interactions between PL cytoplasmic proteins and PDZ proteins to modulate an immune response. In an embodiment, the inhibition or modulation occurs in a hematopoietic cell. In a related embodiment, the inhibition or modulation occurs in an endothelial cell.

As used herein a cytokine receptor has the normal meaning in the art and refers to a membrane protein with an extracellular domain that specifically binds a cytokine. Exemplary PL cytokine receptor proteins include CDW125 (IL5R), CDW128A (IL8RA), and BRL-1. Thus, in one embodiment, the PL proteins of the invention are PL cytokine proteins. In an embodiment, the invention provides methods and reagents, as detailed herein, for inhibiting interactions between PL cytokine proteins and PDZ proteins to modulate an immune response. In an embodiment, the inhibition or modulation occurs in a hematopoietic cell. In a related embodiment, the inhibition or modulation occurs in an endothelial cell.

As used herein, an adaptor protein means a molecule (e.g., protein) that contributes to the formation of a multimolecular complex by binding two or more other biomolecules. The binding of the two or more other molecules by the adaptor molecule/protein generally involves direct molecular contact between the adaptor protein and each of the two or more other molecules. One exemplary PL adaptor protein is LPAP. Thus, in one embodiment, the PL proteins of the invention are PL adaptor proteins. In an embodiment, the invention provides methods and reagents, as detailed herein, for inhibiting interactions between PL adaptor proteins and PDZ proteins to modulate an immune response. In an embodiment, the inhibition or modulation occurs in a hematopoietic cell. In a related embodiment, the inhibition or modulation occurs in an endothelial cell.

In various embodiments, the PL proteins of the invention are characterized by specific C-terminal (i.e., PL domain) amino acid sequences or amino acid motifs, as described elsewhere in this disclosure.

In various embodiments of the invention, the PL proteins of the invention bind a PDZ protein expressed in T lymphocytes, B lymphocytes, or both T and B lymphocytes. In an embodiment, the PL protein binds a PDZ protein expressed in endothelial cells. In various embodiments, the PL proteins and/or the PDZ protein to which it binds are not expressed in the nervous system (e.g., neurons).

In various embodiments of the invention, the PL protein of the invention binds only one PDZ protein listed in TABLE 2. In other embodiments, the PL protein binds 1 to 3, 3 to 5, or more than 5 different PDZ proteins listed in TABLE 2.

In various embodiments of the invention, the PL protein is expressed or up-regulated upon cell activation (e.g., in activated B lymphocytes, T lymphocytes) or upon entry into mitosis (e.g., up-regulation in rapidly proliferating cell populations).

In various embodiments of the invention, the PL protein is (i) a protein that mediates immune cell (e.g., hematopoietic cell) activation or migration, (ii) a protein that does not mediate apoptosis in a cell type, (iii) a protein that is other than a G-protein coupled seven transmembrane helix receptor, (iv) a protein that is G-protein coupled seven transmembrane helix receptor but not a cytokine receptor, or (v) a protein that is not a G-protein coupled seven transmembrane helix receptor and is a cytokine receptor.

6.1 Detection of PDZ Domain-Containing Proteins Expressed in Hematopoietic Cells

As noted *supra*, the present inventors surprisingly discovered that numerous PDZ proteins are expressed in immune system cells, and play a fundamental biological role in modulation of the immune response. PDZ proteins DLG1 and TIAM-1 have been previously described to be in T cells. The present inventors discovered, using a BLAST search of the Human EST database and the experiments described *infra*, that several additional PDZ proteins are present in hematopoietic cells including MPP1, P-DLG, VELI-1, PSD95, syntenin in T cells and CASK, DLG1, DLG2, ZIP KINASE, syntrophin 2, P-dlg, PSD95, and syntenin in B cells.

To determine the full extent of involvement of PDZ proteins in hematopoietic function, the inventors embarked on a systematic investigation of PDZ proteins in T and B cells. A comprehensive list of PDZ domain-containing proteins was retrieved from the Sanger Centre database (Pfam) searching for the keyword, "PDZ". The corresponding cDNA sequences were retrieved from GenBank using the NCBI "entrez" database (hereinafter,

“GenBank PDZ protein cDNA sequences”). The DNA portion encoding PDZ domains was identified by alignment of cDNA and protein sequence using CLUSTALW. Based on the DNA/protein alignment information, primers encompassing the PDZ domains were designed. The expression of certain PDZ-containing proteins in immune cells was detected by polymerase chain reaction (“PCR”) amplification of cDNAs obtained by reverse transcription (“RT”) of immune cell derived RNA (i.e., “RT-PCR”). PCR, RT-PCR and other methods for analysis and manipulation of nucleic acids are well known and are described generally in Sambrook et al., (1989) MOLECULAR CLONING: A LABORATORY MANUAL, 2ND ED., VOLS. 1-3, Cold Spring Harbor Laboratory hereinafter, “Sambrook”); and Ausubel et al., CURRENT PROTOCOLS IN MOLECULAR BIOLOGY, Greene Publishing and Wiley-Interscience, New York (1997), as supplemented through January 1999 (hereinafter “Ausubel”).

In the experiments summarized in TABLE 2, T-cells (Jurkat E6 cell line) and B-cells (MV 4-11 cell line) were tested for expression of specific PDZ domain containing genes by RT-PCR. RNA was prepared using the “trizol” RNA preparation kit (GIBCO-BRL; Cat. # 15596-018) according to the manufacturer’s recommendations. Briefly, $1-5 \times 10^7$ lymphoblasts were harvested by centrifugation at $200 \times g$ for 10 minutes at 20°C . Cells were resuspended in $100 \mu\text{l}$ PBS buffer and 1 ml of TRIZOL reagent was added per 5×10^6 cells. The cells resuspension was mixed and after 5 minutes incubation at room temperature (RT), chloroform was added at 0.2 ml per ml TRIZOL. The resuspension was vigorously shaken and incubated for 3 more minutes at RT. Samples were then centrifuged at $12000 \times g$ for 15 minutes at 4°C , the aqueous phase was recovered and RNA was precipitated with 2-propanol. The precipitate was collected by centrifugation at $12000 \times g$ for 15 minutes at 4°C , washed with 75% ethanol, finally recollected by another spin at $12000 \times g$ for 15 minutes at 4°C , air dried and resuspended in an appropriate volume of DEPC treated water.

RNA concentration and purity were determined by the measurement of 260/280 nm light absorption by the nucleic acid. For cDNA synthesis, the SUPERScript II reverse transcriptase cDNA kit (GIBCO-BRL; Cat. # 18064-014) was used. RNA input per $200 \mu\text{l}$ cDNA reaction sample was $10 \mu\text{g}$. Prior to cDNA synthesis RNA was treated with 1 unit/ μl DNase I in $110 \mu\text{l}$ water at 37°C for 20 minutes. DNase I was then inactivated by a 10 minutes incubation at 70°C . Random primer was used for cDNA priming; $10 \mu\text{l}$ of random hexamer primer ($100 \text{ ng}/\mu\text{l}$) was added, samples were heated to 70°C for 5 minutes and chilled on ice. Subsequently $40 \mu\text{l}$ SUPERScript II “first strand” buffer, $20 \mu\text{l}$ of 0.1 M DDT, $10 \mu\text{l}$ of a 10

mM of mix of deoxynucleotide triphosphates (dATP, dCTP, dGTP, dUTP) and 10 µl of SUPERScript II reverse transcriptase were added and cDNA synthesis was done for 45 minutes at 42°C. Reactions were stopped by a 5 minutes incubation at 95°C and typically, 2-4 µl of such cDNA samples were used for PCR.

5 A portion of the cDNA (typically, 1/5 of a 20 µl reaction) was used for PCR. PCR was conducted using primers designed to amplify specifically PDZ domain-containing regions of PDZ proteins of interest. Oligonucleotide primers were designed to amplify one or more PDZ-encoding domains. The DNA sequences encoding the various PDZ domains of interest were identified by inspection (i.e., conceptual translation of the PDZ protein cDNA sequences obtained from GenBank, followed by alignment with the PDZ domain amino acid sequence). **TABLE 3** shows the PCR primers, the PDZ-encoded domains amplified, and the GenBank accession number of the PDZ-domain containing proteins. To facilitate subsequent cloning of PDZ domains, the PCR primers included endonuclease restriction sequences at their ends to allow ligation with pGEX-3X cloning vector (Pharmacia, GenBank XXI13852) in frame with glutathione-S transferase (GST).

TABLE 3 lists proteins detected in the aforementioned assays. The results showed that PDZ proteins are widely utilized in T and B cells in both lineage specific as well as lineage independent manner. For example INADL2/3 (PDZ dom.), KIAA0316, and 26s subunit p27 were detected in T cells, but not B cells. mCASK, KIAA0559, PTN-4, and X11 beta were detected in B cells, but not T cells. AF6, BAI1 associated prot., Cytohesin bind. Prot., DLG1, DLG5 (pdlg), DVL1, DVL3, GTPase, hypoth. 41.8 kd, KIAA147, KIAA0300, KIAA0303, KIAA0380, KIAA0440, KIAA0545, KIAA0561, LIMK1, LIMK2, LIM domain prot, LIM protein, MINT1, MINT3, MPP1, MPP2, NE-DLG, NOS1, novel serine protease, PTN-3, prIL 16, PSD95, RGS12, serine protease, SYNTENIN, SYNTR 1 alpha, TAX1, TAX2, TAX33, TAX40, Tax43 (SYN, Beta1), TIAM wwp3, and X11 prot. were detected in both T cells and B cells. Similar expression patterns will be apparent from inspection of the Table.

Key: Gene names and corresponding gene products are provided. In some cases, cDNA sequences representing the same gene have several database entries under different accession numbers and names. Accession numbers shown correspond to the gene name used in this description, and numbering of nucleotides and amino acids correlates to the Genbank entry versions specified by the given accession number. Amino acid sequences shown correspond to the cloned DNA portions of PDZ domain containing genes. As is apparent from the primer sequences, in some constructs, the first N-terminal and / or last C-terminal amino acid corresponds to a linker amino acid introduced by the cloning process but is not represented at that position in the corresponding gene. PCR primers were designed such that restriction nuclease recognition sites were generated at the ends of the RT-PCR generated fragments. Therefore, 5' primer sequences do not entirely match with the corresponding cDNA sequences.

GENE SYMBOL	PROTEIN	ACC.#	AMINO ACID SEQUENCE	CLON. SITES	FORWARD PRIMER	REVERSE PRIMER
CASK	Homo sapiens CASK protein	Y17138 GI: 3087817	AA 495 - 584 PDZ domain 1 (of 1) TRVRLVQFQKNTDEPMGITLKMNELNHC IVARIMHGGMIHRQGTSLHVGDEIREING ISVANQTVEQLQKMLREMRGSIITFKIVP SYRTQS	Bam HI / Eco RI	6CAF 5'- TCGGATCCATGT GACCAGAGTTCG G-3' N1471-1494	7CAR 5'- TCGGAATTCAGAC TGAGTCCGGTA- 3' N1761-1738
MPP1	55 Kd erythrocyte membrane protein	M64925 GI: 189785	AA 101 - 186 PDZ domain 1 (of 1) RKVRLIQFEKVTEPFGITLKLNEKQSC TVARILHGGMIHRQGSLSLHVGDEILEING TNVTNHSVDQLQKAMKETKGMISLKVIP NQ	Bam HI / Bam HI	62MPF 5'- GGGATCCGGAAA GTGCGACTCATA C-3' N296-320	63MPR 5'- ACGGATCCGCTGG TTGGGAATTACTT -3' N568-543
LIMK1	human LIM	NM	AA 194 - 291	SMA I	52LIFP	53LIRP

KIAA 0303 (K303)	domain kinase 1	002314 GI: 8051616	PDZ domain 1 (of 1) VTILVSI PASSHGKRLSVSIDPPHPPG CGTEHSHTVRVQGVDPGCMSPDVKNSIH VGDRIEINGTPIRNVPLDEIDLLIQET SRLLQLTLEHD		5'- CTGCCCGGGACC GTACACCTGGTG TCC-3' N570-597	5'- TCGCCCGGGTCAT GCTCGAGGGTC- 3' N874-851
KIAA 0303 (K303)	KIAA 0303 protein	Ab002301 GI: 2224546	AA 652 - 742 PDZ domain 1 (of 1) PHQPIVHSSGKNYGFIRAIRVYVGD DIYTVHHIVMNVEGSPACQAGLKAGDL ITHINGEPVHGLVHTEVIELLLKSGNKV SITTPF	Bam HI / Eco RI	152KIF 5'- CTGGGATCCAC ATCAGCGGATTG TGA-3' N1948-1976	153KIR 5'- TGTGAATTCAAAT GGGTTAGTAGTGA TTG-3' N2237-2209
KIAA 0807 (K807)	KIAA 0807 protein	AB018350 GI: 3882334	AA 635 - 743 PDZ domain 1 (of 1) PIIHRAGKKGFTLRAIRVYMGDSVY TVHHMVHVEDGGPASEAGLRQGLITH VNGEPVHGLVHTEVVELLLKSGNKVAIS TTPL	Bam HI / Eco RI	281KIF 5'- GCAGGATCCCTC CCATCATCATCC AC-3' N1894-1919	282KIR 5'- GATGAATTCTCCA GGGAGTTGTTG- 3' N2155-2179
DLG1	human homolog of Drosophila discs large protein	U13897 GI: 558437	AA 275 - 477 PDZ domains 1-2 (of 3) VNGTDADYEYEITLERGNSGLGFSIAG GTDNPHIGDDSSIFITKIITGGAQAQDG RLRVNDICILQVNEVDVDRVTHSKAVEAL KEAGSIVRLYVRRKPVSEKIMEIKLIK GPKGLGFSIAGGVGNQHIPGDNSIYVTK IIEGGAHKGKGLQIGDKLLAVNNVCLE EVTHSEAVTALKNTSDFVYLKVAKPTSM	Bam HI / Eco RI	1DF 5'- TCGGATCCAGGT TAATGGCTCAGA TG-3' N815-841	2DR 5'- CGGAATTCGGTGC ATAGCCATC-3' N1442-1421

			YNDGYA	Bam HI / Eco RI		
PSD95	human post-synaptic density protein 95	U83192 GI: 3318652	AA 387 - 724 PDZ domains 1-3 (of 3) EGEMEYEEITLERNGSLGFSIAGGTDN PHIGDDPSIFITKIIIPGAAAOQDRLRV NDSILFVNEVDVREVTHSAAVEALKEAG SIVRLYVRRKPPAEKVMIEIKIKGPKG LGFSIAGGVGNQHIPGDNISYVTKIIEG GAAHKDGRQLQIGDKILAVNSVGLEDVMH EDAVAALKNTYDVVYLKVAKPSNAYLSD SYAPPDITTSYSQHLNIEISHSSYLGTD YPTAMTPTSPRRYSPVAKOLLGEEDI PR EPRRIVIHGSGTGLGFNIVGGEDGEGIF ISFILAGGPADLSGELRKQDQILSVNGV DLRNASHEQAAI ALKNAGQTVTIIAQYKPE		8PSF 5'- TCGGATCCTTGA GGGGGAGATGGA -3' N1150-1173	11PSR 5'- TCGGAATTCGCTA TACTCTTCTGG- 3' N2191-2168
NeDLG	Pre-synaptic protein sap102 (neuroendocrine-dlg)	U49089 GI: 1515354	AA 205 - 389 PDZ domains 1-2 (of 3) YEEIVLERNGSLGFSIAGGIDNPHVPD DPGIFITKIIIPGAAAMDGRGLGVNDCVL RVNEVEVSEVHSAVEALKEAGPVVRL VVRRRQPPPETIMEVNLKGPGLGFSI AGGIGNQHIPGDNISYITKIEGGAAQK DGRQLIGDRLLAVNNTNLQDVRHEEAVA SLKNTSDMVYLKVAKPGS	Bam HI / Eco RI	71NEDF 5'- CAGGATCCAATA TGAGGAAATCGT ACTTG-3' N608-635	72NEDR 5'- TTGAATTCGAGGC TGCCCTGGCTTGGC -3' N1186-1161
Syn-trophin alpha 1 gene	Syn-trophin alpha 1 protein	U40571 GI: 1145727	AA 96 - 189 PDZ domain 1 (of 1) QRRRVTVRKADAGGLGISIKGGRENKMP	Bam HI / Eco RI	124SYF 5'- TACGGATCCAGC GGCCGCCGCGTG	125SYR 5'- GTAGAATTCCTGA AATACGGTGAGAC

Table 3
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(SNTa1)			ILISKIFKGLAADQTEALFVGDAILSVN GEDLSSATHDEAVQVLKKTGKEVWLEVK YMKDVSPYFK			AC-3' N279-301	-3' N576-551
TAX-IP 43	human tax interaction protein 43	AF028828 GI: 2613011	AA 15 - 85 PDZ domain 1 (of 1) QKRGVKVLKQELGGLGISIKGGKENKMP ILISKIFKGLAADQTEALYVGDAILSVN GADLRDATHDEAVQAL	Bam HI / Eco RI	97TAF 5'- TCTGGATCCAGA AGCGTGGCGTGA AGG-3' N37-63	98TAR 5'- CGGAATTCAACGC CTGCACCGCCTC- 3' N267-231	
Lim domain protein gene (LDP)	Lim domain protein clp- 36	U90878 GI: 2957144	AA 46 - 88 PDZ domain 1 (of 1) RGMTTQQIDLQGGPWGFRFLVGRKDFEQ PLAISRVTPGSKAAL	Bam HI / Eco RI	146LIF 5'- CCAGGATCCGCG GAATGACCACCC AGC-3' N129-155	147LIR 5'- CATGAATTGCTA GAGCGCCTTGCT T-3' N276-239	
Lim protein gene (LIM)	Human LIM protein	AF061258 GI: 3108092	AA 29 - 112 PDZ domain 1 (of 1) SNYSVSLVGPAPWGFRFLQGGKDFNMPLT ISSLKDGKAAQANVRIGDVVLSIDGIN AQMTHLEAQNKIKGCTGSLNMTLQRAS	Bam HI / Eco RI	182LF 5'- TTAGGATCCTGA GCAAGTACAGTG TGTCAC-3' N86-115	183LR 5'- CTTGAATTCAGCA GATGCTCTTTGCA GAGTC-3' N350-320	
MINT1	human X11 protein	L04953 GI: 340408	AA 717 - 894 PDZ domains 1-2 (of 2) SENCCKDVFIKQKGEILGVVIVESGWGS ILPTVIIANMHGGPAEKSGKLNIGDQI MSINGTSLVGLPLSTCQSIKGLNQSR VKLNIVRCPPVTTLIRRPDLRYQLGFS VQNGIICSLMRGGIAERGGVRVGHRIIE	Eco RI / Eco RI	34MIF 5'- CGGAATTCGGAA AACTGTAAGAT- G-3' N2149-2167	20MR 5'- TCGGAATTCAGCA GCCTGTACATCG- 3' N2690-2666	

X11 beta	Homo sapiens adaptor protein X11-beta	AF047348 GI: 3005559	INGQSVVATPHEKIVHILSNAVGEIHKM TMPAAMYRLLL	Bam HI / Eco RI	133 XF 5'- ACCGATCCACT TCTCAAACTCGG AG-3' N1865-1890	134 XR 5'- AGCGAATTCCT GACCCGTGAGGAG C-3' N2422-2438
KIAA 0440 (K440)	KIAA 0440 protein	AB007900 GI: 2662160	AA 285 - 362 PDZ domain 1 (of 1) SSVEMTLRRNGLQGLGFHVNYEGIVADV EPYGYAWQAGLRQGSRLVEICKVAVATL SHEQMIDLLRTSVTVKVIIPPE	Eco RI / Eco RI	230KIF 5'- AGGGAATTCATC GGTGGAGATGAC TCTGC-3' N843-871	231KIR 5'- CAGAAATTCATGCG GGGGAATGATGAC AAC-3' N1066-1094
KIAA 0545 (K545)	KIAA 0545 protein	AB011117 GI: 3043613	AA 308 - 390 PDZ domain 1 (of 1) SGWETVDMTLRRNGLQGLGFHVKYDGTV AEVEDYGFAWQAGLRQGSRLVEICKVAV VTLTHDQMDIDLLRTSVTVKVIIPPE	Eco RI / Eco RI	293TF 5'- CCGGATCCCGAG GCGAGACCAAGG AGGTG-3' N384-411	294TR 5'- AATGAATTCGAAG GCCCTCTTGGGCT G-3' N672-646
TAX-IP2	human tax interaction protein 2	AF028824 GI: 2613003	AA 54 - 140 PDZ domain 1 (of 1)	Bam HI / Eco RI	197TF 5'- AGGGATCCGCA	198TR 5'- TGTGGAATTCCTT

Table 3
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			RKEVEVFKSEDALGLTITDNGAGYAFIK RIKESVIDHIHLISVGDMEIANGQSL LGCRRHYEVARLLKELPRGRFTTLKLT RK			AGGAGGTGGAGG TGTTTC-3' N154-182	GCGAGGCTCCGTG AGC-3' N429-401
TAX-IP 2- like	human tax interaction 2-like protein	AC005175 GI: 3253116	AA 130 - 221 PDZ domain 1 of 1 IRGETKEVEVTKTEDALGLTITDNGAGY AFIKRIKESIIINRIEAVCVGDSIEAIN DHSIVGCRHYEVAKMLRELPKSQPFTLR LVQPKRAFE	Bam HI / Eco RI	293TF 5'- CCGGATCCCGAG GCGAGACCAAGG AGGTG-3' N384-411	294TR 5'- AATGAATTCGAAG GCCCTCTTGGGCT G-3' N672-646	
TAX-IP 33	tax inter- action protein 33	AF028826 GI: 2613007	AA 73 - 162 PDZ domain 1 (of 1) HSHPRVVELPKTDEGLGFNVMGKEQNS PIYISRIIPGGVAERHGGGLKRGDQLLSV NGVSVEGEHHEKAVELLKAAKDSVKLVV RYTPKVL	Bam HI / Eco RI	92TAF 5'- GTGGATCCACT CCCACCTCGAG TAG-3' N208-234	93TAR 5'- CATGAATTCGAGA ACTTTTGGGTGTA TCGC-3' N497-468	
MPP2	maguk p55 subfamily member 2 (DLG2)	X82895 GI: 939884	AA 185 - 273 PDZ domain 1 (of 1) PVPPDAVRMVGIRKTAGEHLGVTFRVEG GELVIARIILHGGMVAQQGLLHVGDIIKE VNGQPVGSDPRALQELLRNASGSVILKI LPNYQ	Bam HI / Eco RI	142MF 5'- TCAGGATCCAGC CTGTACCTCCCG ATGC-3' N542-569	143MR 5'- ATGGAATTCCTGG TAGTTGGGCAGGA TC-3' N828-801	
MINT3	human MINT3	AF029110 GI: 3169808	AA 11 - 52 PDZ domain 1 (of 1) PVTTAIIHRPHAREQLGFCVEDGIVRPR	Bam HI / Eco RI	188MF 5'- ACTGGATCCCGG TCACCACCGCCA	189MR 5'- CTCGAATTCCTGTG CTCAGGGCCGCCCC	

			PLAPGWGGRALST			TCATC-3'	TA-3'
						N23-51	N165-138
TIP-1	Homo sapiens Tax interaction protein 1	AF028823 GI: 2613001	AA 14 - 117 PDZ domain 1 (of 1) QRVEIHKLRQGENLILGFSIGGIDQDP SQNPFSDEKTDKGIYVTRVSEGGPAEIA GLQIGDKIMQVNGWDMTMTVTHDQARKRL TKRSEEVVRLLVTRQSLQK	Bam HI / Eco RI	86TAF 5'- CAGGGATCCAAA GAGTTGAAATTC ACAAGC-3' N10-39	87TAR 5'- ACGGAATTCGCA GCGACTGCCGCGT C-3' N305-331	
PTN-4	protein- tyrosine phosphatase meg1	M68941 GI: 190747	AA 774 - 862 PDZ domain 1 (of 1) LIRMKPDENGFRGFNVKGGYDQKMPVIV SRVAPGTPADLCVPRINEGDQVVLINGR DIAETHDQVVLFIKASCERHSGELMLL VRPNA	Bam HI / Eco RI	247PTF 5'- ATCGGATCCTAA TCAGAATGAAAC CTG-3' N2312-2338	248PTR 5'- ATCGAATTCAGCA TTAGGTCGAACTA G-3' N2595-2569	
prIL16	putative interleukin 16 precursor	S81601 GI: 1478492	AA 170 - 383 PDZ domain 1-2 (of 2) IHVTILHKEEGAGLGFSLAGGADLENKV ITVHRVFPNGLASQEGTIQKNEVLIN GKSLKGTTHDALAILRQAREPRQAVIV TRKLTPEAMPDLNSSTDASAASAASDV SVESTAEATVCTVLEKMSAGLGFSLEG GKSLHGDKPLTINRIFKGAASEQSETV QPGDEILQLGGTAMQGLTRFEAWNIIKA LPDGPVTIVIRKSLQSK	Bam HI / Eco RI	75PRF 5'- ACGGATCCATG TCACCATCTTAC AC-3' N503-528	76PRR 5'- GTGAATTCCTTGG ACTGGAGGCTTTT TC-3' N1157-1129	
Cyto-hesin	Cytohesin	AF08836	AA 85 - 76	Bam HI /	235CYF	236CYR	

Table 3
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binding protein gene (CBP)	binding protein HE	GI: 3192908	PDZ domain 1 (of 1) QRKLVTVREKQDNETFGFEIQSYRPQNQ ACSEMFTLICKIQEDSPAHCAGLQAGD VLANINGVSTEGFTYKQVVDLIRSSGNL LTITETLNG	Eco RI	5'- CCTGGATCCAAA GAAAGCTTGTTA CTGTG-3' N246-274	5'- TCAGAATTCATT AAGAGTCTCTATC -3' N535-510
KIAA 0751 (41.8)	Hypoth. 41.8 kD protein	AF007156 GI: 2852637	AA 4 - 85 PDZ domain 1 (of 1) RDSGAMGLKVVGGKWTESGRLCFAITK VKKGLADTVGHLRPPGDEVLEWNGRLIQ GATFEVYNIILESKEPQVELVVS	Bam HI / Eco RI	145HF 5'- GTGGATCCGAG ATTGAGGAGCAA TGC-3' N4-30	146HR 5'- CTGGAATTCGCCT TGAAGCTACAAGT TC-3' N267-240
KIAA 0559 (K559)	KIAA 0559 protein	AB011131 GI: 3043641	AA 766 - 870 PDZ domain 1 (of 1) HYIFPHARIKITRDSKDHVSGNGLGIR IVGKEIPGHSGEIGAYIAKILPGGSAE QTGKLMGMOVLEWNGIPLTSKTYEEVQ SIISQQSGEAEICVRLDLNML	Bam HI / Eco RI	130KIF 5'- AAAGGATCCACT ACATCTTTCCTC ACG-3' N2290-2312	131KIR 5'- TCACAATTGGATA GCATATTGAGGTC CAG-3' N2623-2595
AF6	af-6 protein	U02478 GI: 430993	AA 985 - 1077 PDZ domain 1 (of 1) LRKEPEIITVTLKKQNGMGLSIVAAGGA GQDKLGIYVKSVVKGGAADVDRGLAAGD QLLSVDGRSLVGLSQERAAELMTTSSV VTLEVAKQG	Bam HI / Eco RI	66AFF 5'- TCGGATCCTGAG GAAAGAACCTGA A-3' N2946-2970	67AFR 5'- TAGAATTCACCT GCTTTGCTACTTC -3' N3239-3214
PICK1	Novel human mRNA similar	AL049654 GI:	AA 16 - AA 105	Bam HI / Eco RI	287PIF	288PIR

	to mouse gene	4678411	PDZ domain 1 (of 1) PTVPGKVTLOKDAQNLIIGISIGGAQYC PCLYIVQVFDNTPAALDGTVAAGDEITG VNGRSIKGKTKVEVAKMIQEVKGE VTIHYNKLQE		5'- TCGGGATCCCGA CTGTGCCTGGGA AG-3' N268-N293 64RGR	5'- CTTGAATTCCTCC TGCAGCTTCTTGT TGTAAG-3' N527-N554
RGS12	human regulator of G-protein signal-ling 12	AF035152 GI: 3290015	AA 35 - 103 PDZ domain 1 (of 1) PPRVRSEVGRAGYFTLSGQAPCVL SCVMRGSPADFVGLRAGDQILAVNEINV KKASHEDVVKLIG	Bam HI / Eco RI	5'- TGGGATCCCGCC CCCAAGGTGCG GAG-3' N93-119 238PDF	5'- AGGAATTCCTCAAT TAATTTCACTAC- 3' N316-291 239PDR
PDZK1	Homo sapiens PDZ domain contain-ing protein (PDZK1)	AF012281 GI: 2944188	AA 134 - 457 PDZ domains 2 - 4 (of 4) RLCYLVKEGGSYFSLKTVQKKGVYMT DITPQGVAMRAGVLADDDHLIEVNGENVE DASHEKWEKVKKSGSRVWFLVLDKETD KRHVEQKIQFKRETASLKLPHQPRIVE MKKGSNGYGFYLRAGSEKQGIKIDIS GSPAEEAGLKNNDLVVAVNGESVETLDH DSVEMIRKGGDQTSLLVVDKETDNMYR LAHFPFLYYQSQELPNSVKEAPATP TSLEVSSPPDTTEEVDPKLCRLAKGE NGYGFHLNAIRGLPGSFIKEVQKGGPAD LAGLEDEDVILIEVNGVNVLDPEYKVVVD RIQSSGKNVTLLVCGK	Bam HI / Eco RI	5'- CCGGATCCGGCT CTGCTATCTCGT GAA-3' N 426 - 452 N 1385 - 1412	5'- TAGGAATTCCTTC CTCAGACTAGAAG TG-3' N 1385 - 1412
KIAA 0316 (K316)	KIAA 0316 protein	AB002314 GI: 6683123	AA 197 - 284 PDZ domain 1 (of 1) IPPAPRKVEMRRDPVLGFGFVAGSEKPV	Bam HI / Eco RI	158KIF 5'- AAAGGATCCCTC CGGCTCCTCGGA	159KIR 5'- TTAGAATTCGTAT TTGGGAGAGGGT

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			VVRSVTPGSPSEGLIPGDQIVMINDEP VSAAPRRVIDLVRSCKESILLTVIQPY PSPK			AG-3' N586-611	AAG-3' N866-839
DLG5	Human discs large protein p-dlg	U61843 GI: 3650451	AA 99 - 338 PDZ domains 2 (of 2) PYVEEPRHVQKVGSEPLGISIVSGEKG GIYVSKVTVGSIAHQAGLEYGDQLLEFN GINLRSATEQOARLIIGQQCDTITILAQ YNPHVQLSSHSRSSHLDPAETHSTLQ GSGTTTPEHPSVIDPLMEQDEGPSTPPA KQSSSRIAGDANKKLEPRVVFIIKSQ ELGVHLGGNLHGVFAEVEDDSPA DGLVPGDLILEYGSIDVRNKTVEEVYVE MLKPRDGVRLKVQYRPE	Bam HI / Eco RI	81PDLGF 5'- ATAGATCCCTT ATGTGGAGGAGC CAC-3' N645-N671	82PDLGR 5'- TTGAATTCCTCAG GGCGGTACTGCAC CTTC-3' N1356-N1385	
Mouse Syntenin gene (SYNT)	Mus musculus Syntenin	AF077527 GI: 3342559	AA 67 - 241 REIKQIREVILCKDQDGKIGLRKSID NGIFVQLVQANSPASLVGLRFGDQVLQI NGENCAGWSSDKAHKVLKQAFGEKITMT IRDRPFERTVIMHKDSSGHVGFIFKSGK ITSIVKSSAARNGLLTDHHICEINGQN VIGLKDAQIADILSTAGTVVT ITIMPTTFIFEHIIRMAPSM	Bam HI / Eco RI	14SF 5'- TCGGATCCTTGA AATTAAGCAAGG GAT-3' N363-N390	15SR 5'- TCGGAATTCATGC CTGGAGCCATCC- 3' N896-N920	
WWP3	Homo sapiens membrane associated guanylate kinase 1	U80754 GI: 2695619	AA 314 - 576 PDZ domains 1-2 (of 2) PSELKGKFIHTKLRKSSRGFGFTVVGDD EPDEFLLQIKSLVLDGPAALDGKMETGDV	Bam HI / Eco RI	164WWF 5'- CACGGATCCCTT CTGAGTTGAAAG GC-3'	165WWR 5'- CTTGAATTCCTGGC AGCCCTCTCTGTT GC-3'	

	(MAGI-1)			IVSVNDTCVLGHTHAQVVKIFQSIPIGA. SVDLELCRGYPLPFDPPDNTSLVTSVA ILDKPIIVNGQETYDSPASHSSKTGKV NGMKDARPPSSPADVASNSSH GYPNDTVSLASSIATQPELITVHIVKGP MGFGFTIADSPGGGQGVKQIVDSRCCR GLKEGDLIVEVNKKVQALTNQVVDML VECPKGSSEVTLVQRGGILP			N932-N957	N1710-N1737
TAX-IP 40	human tax inter-action protein 40	AF028827 GI: 2613009	AA 35 - 137 PDZ domain 1 (of 1)	PDZ domain 1 (of 1)	Bam HI / Eco RI	136TF 5'- ACGGGATCCTAC TGCCTGAGACCC ACC-3' N97-123	137TR 5'- ACGGAATCCGCT GGTGGCGGGCTT GAC-3' N421-393	
KIAA 0858 (K858)	KIAA 0858 protein	AB020665 GI: 4240204	AA 66 - 159 PDZ domain 1 (of 1)	PDZ domain 1 (of 1)	Bgl II / Eco RI	278KIF 5'- AGGAGATCTTCA GTGATATGAGAA TC-3' N190-N215	279KIR 5'- CTTGAATTCAGGT GAACGAGCCTTTC -3' N460-N485	
TIAM1	T- lymphoma invasion and metastasis inducing protein 1	NM_ 003253 GI: 4507500	AA 1001 - 1088 PDZ domain 1 (of 1)	PDZ domain 1 (of 1)	Bam HI / Eco RI	39TF 5'- TCGGATCCACAG CATCCACATTGA G-3' N2995-3019	40TR 5'- TCGGAATTCCTCC AGCTCGGGGT-3' N3275-3253	
Connector	Homo sapiens	AF100153	AA 193 - 300	AA 193 - 300	Bam HI /	296CF	297CR	

Enhancer gene (ConEn)	connector enhancer of KSR-like protein CNK1	GI: 3930780	PDZ domain 1 (of 1) LEQAVLEQVQLDSPGLGLEIHTSNCQH FVSQVDTQVPTDSRLQIQPGDEVVQINE QVVVGWPRKNMVRLLREPAGLSL VLKKIPIP	Eco RI	5'- AGGGATCCTGG AACAGAGGCCG TGCTC-3' N605-N633	5'- GGGAATCCGGTA TCGGATCTTCCT TC-3' N858-N884
Serine protease (SPsht)	Homo sapiens serine protease (omi)	AF020760 GI: 2738914	AA 421 - 506 Splice variant: void of AA 444 - 465 (ref. to GI: 2738914) PDZ domain 1 (of 1) SSSGISGSQRRYIGVMMLTLSPSAGLRP GDVILAIGEVMQVNAEDVYEAVRTQSE	Eco RI / Eco RI	191SF 5'- GAAGAATTCCTC CTCCGAATCAG TG-3' N1501-N1526	192SR 5'- TGCGAATTCGGAT TGGGTTCGAACAG CTTC-3' N1774- N1803
DVL1	human dishevelled segment polarity protein homolog	AF006011 GI: 2291005	AA 248 - 340 PDZ domain 1 (of 1) LNIVTVTLNMRHHFLGISIVGQSNDRG DGGIYIGSIMKGGAVADGRIEFGDMLL QVNDVNFENMSNDDAVRVLREIVSQTGP ISLTVAKCW	Bam HI / Eco RI	1 st PCR: 55DVISF 5'- TCATCCAGACTC ATCCGGAAG-3' N652-673 2 nd PCR, nested: 37DVF 5'- TCGGATCCAAAC GGTCACTCTCAA	1 st PCR: 56DVISR 5'- GCTCATGTCACTC TTCACCG-3' N1195-1174 2 nd PCR, nested: 38DVR 5'- TCGGAATTCCTCAG CACTTGGCTACAG

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						C-3'	-3'
Novel serine protease (NSP)	Homo sapiens novel serine protease protein (PRSS11)	Y07921 GI: 1621243	AA 107 - 204 PDZ domain 1 (of 2) IRQAKGKAITKKYIGIRMSLTSSKAK ELKDRHRDFPDVISGAYIIIEVIPDPFAE AGGLKENDVIISINGQSVVSANDVSDVI KRESTINMVRNGN	Bam HI / Eco RI	194NSP 5'- CCCGATCCGAC AGCCAAAGGAA AAGC-3' N1138-N1165	195NSR 5'- GATGAATTCATTA CCCCTGCGGACCA CCATG-3' N1415- N1445	N1029-N1004
Guanin Change Factor gene (GEF)	Homo sapiens PDZ domain containing guanine nucleotide exchange factor 1	AF117947 GI: 6650765	AA 343 - 450 PDZ domain 1 (of 1) CSVMIFEVVEQAGAILLEDGQELDSWV ILNGTVEISHPDGKVENLFMGNSFGITP TLDKQYMHGIVRTKVDDCQFVCIAQQDY WRILNHVEKNTHKVEEGEIVMVH	Bgl II Eco RI	275GF 5'- GAGAGATCTGCT CAGTGATGATTT TTG-3' N1088-N1114	276GR 5'- CCGGAATTCATGT ACCATAACAATTT C-3' N1402- N1428	
KIAA 0902 (K902)	KIAA 0902 protein	AB020715 GI: 4240304	AA 214 - 301 PDZ domain 1 (of 1) ILNEMIAPMVRVNYGOSTDINAFVGAVS LSCSDSGLWAVEGKNKLVCSGLLQASKS NLISGSVMYIEEKTTKYTGNTKMYEV VYQIG	Bam HI / Eco RI	290KIF 5'- AGAGGATCCTCA ATGAAATGATTTG C-3' N633-N657	291KIR 5'- TCTGAATTCCAAT TTGGTAGACCACT TC-3' N884 -N991	
KIAA 0561 (K561)	KIAA 0561 protein	AB011133 GI: 3043645	AA 948 - 1038 PDZ domain 1 (of 1)	Bam HI / Eco RI	161KIF 5'- CCTGGATCCCCC	162KIR 5'- GAGGAATTCCTCA	

				PPSLSTALARSTASACGRSASTWVIATS TLCTTSSGVWRTEAPRRRACGLGTSSP TSTGSCWGWCCTWTWSGCCZRAATRYPC GPQPWR			CATCGTTATCCA CAGC-3' N2836-2863	GGGCTGTGTGTCGG -3' N3120-3095
NOS1	human neuronal nitric oxide synthase	U17327 GI: 642525	AA 239 - 329 PDZ domain 1 (of 1) IQPNVISVRLFKRKVGGGLGFLVKERVSK PPVIISDLIRGGAEEQSGLIQAGDILIA VNGRPLVDLSYDSALEVLRLGIASETHVV LILRGP	Bam HI / Eco RI	155NOF 5'- AGCGATCCAGC CCAATGTCAATTT C-3' N711-733	156NOR 5'- GAAGAATTCAGGG CCCCTCAGAATG- 3' N994-970		

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*Note concerning TABLE 3

In several cases, sequence analysis of the PDZ clones revealed differences to the DNA and/or protein sequence as published in the databases, summarized in TABLE 3A.

TABLE 3A

GENE	GENBANK ENTRY***	ACTUAL CONSTRUCT
AF6	N 3060: C	N 3060: T *
DLG1	N 1021: A, = AA 340: Gln	N 1021: G, = AA 340: Arg
Lim dom.	N 202: G N 203: C, = AA 68: Arg	N 202: C * N 203: G, = AA 68: Gly
LIMK1	N 855: C, = AA 285: Leu	N 855: A, = AA 285: Ile
MINT1	N 2386: G, = AA 796: Glu**	N 2386: A, = AA 796: Lys**
NE-DLG	N 713: T N 766: G, = AA 255: Gly N 803: G, = AA 267: Glu N 861: G, = AA 287: Val	N 713: C* N 766: A, = AA 255: Glu N 803: C, = AA 267: Asp N 861: A, = AA 287: Met
TIAM1	N 3224: A	N 3224: G*
MPP2	N 812 = A; AA = Asn	N 812 = G; AA= Ser
TIP-1	N 196 = T; AA = Ile	N 196 = G; AA= Ser

(*) = silent mutation, does not effect the AA sequence;

(**) = MINT1 is the same as X11a. The database entry for X11a shows the same sequence as our actual construct with regard to N 2386 of the MINT1 GenBank entry.

(***)= Nucleotide ("N") and amino acid ("AA") annotations correspond to the numbering as found in the GenBank files (for accession no., see Table 3).

6.2 Assays for Detection of Interactions Between PDZ-Domain Polypeptides and Candidate PDZ Ligand proteins (PL proteins)

Two complementary assays, termed "A" and "G," were developed to detect binding between a PDZ-domain polypeptide and candidate PDZ ligand. In each of the two different assays, binding is detected between a peptide having a sequence corresponding to the C-terminus of a protein anticipated to bind to one or more PDZ domains (i.e. a candidate PL peptide) and a PDZ-domain polypeptide (typically a fusion protein containing a PDZ domain). In the "A" assay, the candidate PL peptide is immobilized and binding of a soluble PDZ-domain polypeptide to the immobilized peptide is detected (the "A" assay is named for the fact that in one embodiment an avidin surface is used to immobilize the peptide). In the "G" assay, the PDZ-domain polypeptide is immobilized and binding of a soluble PL peptide is detected (The "G" assay is named for the fact that in one embodiment a GST-binding surface is used to immobilize the PDZ-domain polypeptide). Preferred embodiments of these assays are described in detail *infra*. However, it will be appreciated by ordinarily skilled practitioners that

these assays can be modified in numerous ways while remaining useful for the purposes of the present invention.

6.2.1 Production of Fusion Proteins Containing PDZ-Domains

GST-PDZ domain fusion proteins were prepared for use in the assays of the invention. PCR products containing PDZ encoding domains (as described in §6.1 *supra*) were subcloned into an expression vector to permit expression of fusion proteins containing a PDZ domain and a heterologous domain (i.e., a glutathione-S transferase sequence, "GST"). PCR products (i.e., DNA fragments) representing PDZ domain encoding DNA was extracted from agarose gels using the "sephaglas" gel extraction system (Pharmacia) according to the manufacturer's recommendations.

As noted *supra*, PCR primers were designed to include endonuclease restriction sites to facilitate ligation of PCR fragments into a GST gene fusion vector (pGEX-3X; Pharmacia, GenBank accession no. XXU13852) in-frame with the glutathione-S transferase coding sequence. This vector contains a IPTG inducible lacZ promoter. The pGEX-3X vector was linearized using *Bam* HI and *Eco* RI or, in some cases, *Eco* RI or *Sma* I, as shown in TABLE 3, and dephosphorylated. For most cloning approaches, double digestion with *Bam* HI and *Eco* RI was performed, so that the ends of the PCR fragments to clone were *Bam* HI and *Eco* RI. In some cases, restriction endonuclease combinations used were *Bgl* II and *Eco* RI, *Bam* HI and *Mfe* I, or *Eco* RI only, *Sma* I only, or *Bam*HI only (see TABLE 3). When more than one PDZ domain was cloned, the DNA portion cloned represents the PDZ domains and the cDNA portion located between individual domains. Precise locations of cloned fragments used in the assays are indicated in TABLE 3. DNA linker sequences between the GST portion and the PDZ domain containing DNA portion vary slightly, dependent on which of the above described cloning sites and approaches were used. As a consequence, the amino acid sequence of the GST-PDZ fusion protein varies in the linker region between GST and PDZ domain. Protein linkers sequences corresponding to different cloning sites/approaches are shown below. Linker sequences (vector DNA encoded) are bold, PDZ domain containing gene derived sequences are in italics.

- 1) **GST—BamHI/BamHI—** *PDZ domain insert*
Gly—Ile— *PDZ domain insert*
- 2) **GST—BamHI/BglII—** *PDZ domain insert*
Gly—Ile— *PDZ domain insert*

- 3) **GST—EcoRI/EcoI—PDZ domain insert**
Gly—Ile—Pro—Gly—Asn—PDZ domain insert (SEQ ID NO: 360)
- 5 4) **GST—SmaI/SmaI—PDZ domain insert**
Gly—Ile—Pro—PDZ domain insert

The PDZ-encoding PCR fragment and linearized pGEX-3X vector were ethanol precipitated and resuspended in 10 ul standard ligation buffer. Ligation was performed for 4-10 hours at 7°C using T4 DNA ligase. It will be understood that some of the resulting constructs include very short linker sequences and that, when multiple PDZ domains were cloned, the constructs included some DNA located between individual PDZ domains.

The ligation products were transformed in DH5α or BL-21 *E.coli* bacteria strains. Colonies were screened for presence and identity of the cloned PDZ domain containing DNA as well as for correct fusion with the glutathione S-transferase encoding DNA portion by PCR and by sequence analysis. Positive clones were tested in a small scale assay for expression of the GST/PDZ domain fusion protein and, if expressing, these clones were subsequently grown up for large scale preparations of GST/PDZ fusion protein.

GST-PDZ domain fusion protein was overexpressed following addition of IPTG to the culture medium and purified. Detailed procedure of small scale and large scale fusion protein expression and purification are described in "GST Gene Fusion System" (second edition, revision 2; published by Pharmacia). In brief, a small culture (3-5mls) containing a bacterial strain (DH5α, BL21 or JM109) with the fusion protein construct was grown overnight in LB-media at 37°C with the appropriate antibiotic selection (100ug/ml ampicillin; a.k.a. LB-amp). The overnight culture was poured into a fresh preparation of LB-amp (typically 250-500mls) and grown until the optical density (OD) of the culture was between 0.5 and 0.9 (approximately 2.5 hours). IPTG (isopropyl β-D-thiogalactopyranoside) was added to a final concentration of 1.0mM to induce production of GST fusion protein, and culture was grown an additional 1.5-2.5 hours. Bacteria were collect by centrifugation (4500 g) and resuspended in Buffer A- (50mM Tris, pH 8.0, 50mM dextrose, 1mM EDTA, 200uM phenylmethylsulfonylfluoride). An equal volume of Buffer A+ (Buffer A-, 4mg/ml lysozyme) was added and incubated on ice for 3 min to lyse bacteria. An equal volume of Buffer B (10mM Tris, pH 8.0, 50mM KCl, 1mM EDTA, 0.5% Tween-20, 0.5% NP40 (a.k.a. IGEPAL CA-630), 200uM phenylmethylsulfonylfluoride) was added and incubated for an additional 20

min. The bacterial cell lysate was centrifuged (x20,000g), and supernatant was added to glutathione Sepharose 4B (Pharmacia, cat no. 17-0765-01) previously swelled (rehydrated) in 1X phosphate-buffered saline (PBS). The supernatant-Sepharose slurry was poured into a column and washed with at least 20 bed volumes of 1X PBS. GST fusion protein was eluted
5 off the glutathione sepharose by applying 0.5-1.0 ml aliquots of 5mM glutathione and collected as separate fractions. Concentrations of fractions were determined using BioRad Protein Assay (cat no. 500-0006) according to manufacturer's specifications. Those fractions containing the highest concentration of fusion protein were pooled and dialyzed against 1X PBS/35% glycerol. Fusion proteins were assayed for size and quality by SDS gel electrophoresis (PAGE) as
10 described in "Sambrook." Fusion protein aliquots were stored at minus 80°C and at minus 20°C.

6.2.2 Identification of Candidate PL Proteins and Synthesis of Peptides

In some non-hematopoietic cells (e.g., neurons, epithelial cells), certain PDZ domains are known to be bound by the C-terminal residues of PDZ-binding proteins. To
15 identify PL proteins that function in hematopoietic and endothelial cells, cell surface receptor proteins were identified and peptides having the sequence corresponding to the C-terminus of each protein were synthesized. TABLE 4 lists these proteins, and provides corresponding C-terminal sequences and GenBank accession numbers. "Clasp 1" is described in WO 00/20434 (published 13 April 2000). "Clasp 2" and "Clasp 4" are described in copending applications
20 USSN 09/547276, 60/196527, 60/240,503, 09/687837 and PCT/US00/10158 and have the C-terminal sequences shown in Table 4.

Synthetic peptides of defined sequence (e.g., corresponding to the carboxyl-termini of the indicated proteins) can be synthesized by any standard resin-based method (see, e.g., U. S. Pat. No. 4,108,846; see also, Caruthers et al., 1980, *Nucleic Acids Res. Symp. Ser.*,
25 215-223; Horn et al., 1980, *Nucleic Acids Res. Symp. Ser.*, 225-232; Roberge, et al., 1995, *Science* 269:202). The peptides used in the assays described herein were prepared by the Fmoc (see, e.g., Guy and Fields, 1997, *Meth. Enz.* 289:67-83; Wellings and Atherton, 1997, *Meth. Enz.* 289:44-67). In some cases (e.g., for use in the A and G assays of the invention), peptides were labeled with biotin at the amino-terminus by reaction with a four-fold excess of
30 biotin methyl ester in dimethylsulfoxide with a catalytic amount of base. The peptides were

cleaved from the resin using a halide containing acid (e.g. trifluoroacetic acid) in the presence of appropriate antioxidants (e.g. ethanedithiol) and excess solvent lyophilized.

Following lyophilization, peptides can be redissolved and purified by reverse phase high performance liquid chromatography (HPLC). One appropriate HPLC solvent system involves a Vydac C-18 semi-preparative column running at 5 mL per minute with increasing quantities of acetonitrile plus 0.1% trifluoroacetic acid in a base solvent of water plus 0.1% trifluoroacetic acid. After HPLC purification, the identities of the peptides are confirmed by MALDI cation-mode mass spectrometry. As noted, exemplary biotinylated peptides are provided in TABLE 4.

6.2.3 Detecting PDZ-PL Interactions

Based on the determination that immune system cells contain both many PDZ proteins and similarly many candidate PL proteins, it was apparent to the inventors that characterization of the specific PDZ-PL interactions among these proteins would require reliable and rapid assays for such interactions. A variety of assay formats known in the art can be used to select ligands that are specifically reactive with a particular protein. For example, solid-phase ELISA immunoassays, immunoprecipitation, Biacore, and Western blot assays can be used to identify peptides that specifically bind PDZ-domain polypeptides. As discussed *supra*, two different, complementary assays were developed to detect PDZ-PL interactions. In each, one binding partner of a PDZ-PL pair is immobilized, and the ability of the second binding partner to bind is determined. These assays, which are described *infra*, can be readily used to screen for hundreds to thousand of potential PDZ-ligand interactions in a few hours.

Thus these assays can be used to identify yet more novel PDZ-PL interactions in hematopoietic cells. In addition, they can be used to identify antagonists of PDZ-PL interactions (see *infra*).

In various embodiments, fusion protein are used in the assays and devices of the invention. Methods for constructing and expressing fusion proteins are well known. Fusion proteins generally are described in Ausubel et al., *supra*, Kroll et al., 1993, DNA Cell. Biol. 12:441, and Imai et al., 1997, *Cell* 91:521-30. Usually, the fusion protein includes a domain to facilitate immobilization of the protein to a solid substrate ("an immobilization domain"). Often, the immobilization domain includes an epitope tag (i.e., a sequence recognized by a antibody, typically a monoclonal antibody) such as polyhistidine (Bush et al, 1991, *J. Biol Chem* 266:13811-14), SEAP (Berger et al, 1988, *Gene* 66:1-10), or M1 and M2 flag (see, e.g, U.S.

Pat. Nos. 5,011,912; 4,851,341; 4,703,004; 4,782,137). In an embodiment, the immobilization domain is a GST coding region. It will be recognized that, in addition to the PDZ-domain and the particular residues bound by an immobilized antibody, protein A, or otherwise contacted with the surface, the protein (e.g., fusion protein), will contain additional residues. In some
5 embodiments these are residues naturally associated with the PDZ-domain (i.e., in a particular PDZ-protein) but they may include residues of synthetic (e.g., poly(alanine)) or heterologous origin (e.g., spacers of, e.g., between 10 and 300 residues).

PDZ domain-containing polypeptide used in the methods of the invention (e.g., PDZ fusion proteins) of the invention are typically made by (1) constructing a vector (e.g.,
10 plasmid, phage or phagemid) comprising a polynucleotide sequence encoding the desired polypeptide, (2) introducing the vector into an suitable expression system (e.g., a prokaryotic, insect, mammalian, or cell free expression system), (3) expressing the fusion protein and (4) optionally purifying the fusion protein.

(1) In one embodiment, expression of the protein comprises inserting the
15 coding sequence into an appropriate expression vector (i.e., a vector that contains the necessary elements for the transcription and translation of the inserted coding sequence required for the expression system employed, e.g., control elements including enhancers, promoters, transcription terminators, origins of replication, a suitable initiation codon (e.g., methionine), open reading frame, and translational regulatory signals (e.g., a ribosome binding site, a
20 termination codon and a polyadenylation sequence. Depending on the vector system and host utilized, any number of suitable transcription and translation elements, including constitutive and inducible promoters, can be used.

The coding sequence of the fusion protein includes a PDZ domain and an immobilization domain as described elsewhere herein. Polynucleotides encoding the amino
25 acid sequence for each domain can be obtained in a variety of ways known in the art; typically the polynucleotides are obtained by PCR amplification of cloned plasmids, cDNA libraries, and cDNA generated by reverse transcription of RNA, using primers designed based on sequences determined by the practitioner or, more often, publicly available (e.g., through GenBank). The primers include linker regions (e.g., sequences including restriction sites) to facilitate cloning
30 and manipulation in production of the fusion construct. The polynucleotides corresponding to the PDZ and immobilization regions are joined in-frame to produce the fusion protein-encoding sequence.

The fusion proteins of the invention may be expressed as secreted proteins (e.g., by including the signal sequence encoding DNA in the fusion gene; see, e.g., Lui et al, 1993, *PNAS USA*, 90:8957-61) or as nonsecreted proteins.

In some embodiments, the PDZ-containing proteins are immobilized on a solid surface. The substrate to which the polypeptide is bound may in any of a variety of forms, e.g., a microtiter dish, a test tube, a dipstick, a microcentrifuge tube, a bead, a spinnable disk, and the like. Suitable materials include glass, plastic (e.g., polyethylene, PVC, polypropylene, polystyrene, and the like), protein, paper, carbohydrate, lipid monolayer or supported lipid bilayer, and other solid supports. Other materials that may be employed include ceramics, metals, metalloids, semiconductive materials, cements and the like.

In some embodiments, the fusion proteins are organized as an array. The term "array," as used herein, refers to an ordered arrangement of immobilized fusion proteins, in which particular different fusion proteins (i.e., having different PDZ domains) are located at different predetermined sites on the substrate. Because the location of particular fusion proteins on the array is known, binding at that location can be correlated with binding to the PDZ domain situated at that location. Immobilization of fusion proteins on beads (individually or in groups) is another particularly useful approach. In one embodiment, individual fusion proteins are immobilized on beads. In one embodiment, mixtures of distinguishable beads are used. Distinguishable beads are beads that can be separated from each other on the basis of a property such as size, magnetic property, color (e.g., using FACS) or affinity tag (e.g., a bead coated with protein A can be separated from a bead not coated with protein A by using IgG affinity methods). Binding to particular PDZ domain may be determined; similarly, the effect of test compounds (i.e., agonists and antagonists of binding) may be determined.

Methods for immobilizing proteins are known, and include covalent and non-covalent methods. One suitable immobilization method is antibody-mediated immobilization. According to this method, an antibody specific for the sequence of an "immobilization domain" of the PDZ-domain containing protein is itself immobilized on the substrate (e.g., by adsorption). One advantage of this approach is that a single antibody may be adhered to the substrate and used for immobilization of a number of polypeptides (sharing the same immobilization domain). For example, an immobilization domain consisting of poly-histidine (Bush et al, 1991, *J. Biol Chem* 266:13811-14) can be bound by an anti-histidine monoclonal antibody (R&D Systems, Minneapolis, MN); an immobilization domain consisting of secreted

alkaline phosphatase ("SEAP") (Berger et al, 1988, *Gene* 66:1-10) can be bound by anti-SEAP (Sigma Chemical Company, St. Louis, MO); an immobilization domain consisting of a FLAG epitope can be bound by anti-FLAG. Other ligand-antiligand immobilization methods are also suitable (e.g., an immobilization domain consisting of protein A sequences (Harlow and Lane, 1988, *Antibodies A Laboratory Manual*, Cold Spring Harbor Laboratory; Sigma Chemical Co., St. Louis, MO) can be bound by IgG; and an immobilization domain consisting of strepavidin can be bound by biotin (Harlow & Lane, *supra*; Sigma Chemical Co., St. Louis, MO). In a preferred embodiment, the immobilization domain is a GST moiety, as described herein.

When antibody-mediated immobilization methods are used, glass and plastic are especially useful substrates. The substrates may be printed with a hydrophobic (e.g., Teflon) mask to form wells. Preprinted glass slides with 3, 10 and 21 wells per 14.5 cm² slide "working area" are available from, e.g., SPI Supplies, West Chester, PA; also see U.S. Pat. No. 4,011,350). In certain applications, a large format (12.4 cm x 8.3 cm) glass slide is printed in a 96 well format is used; this format facilitates the use of automated liquid handling equipment and utilization of 96 well format plate readers of various types (fluorescent, colorimetric, scintillation). However, higher densities may be used (e.g., more than 10 or 100 polypeptides per cm²). See, e.g., MacBeath et al, 2000, *Science* 289:1760-63.

Typically, antibodies are bound to substrates (e.g., glass substrates) by adsorption. Suitable adsorption conditions are well known in the art and include incubation of 0.5-50ug/ml (e.g., 10 ug/ml) mAb in buffer (e.g., PBS, or 50 to 300 mM Tris, MOPS, HEPES, PIPES, acetate buffers, pHs 6.5 to 8, at 4°C) to 37°C and from 1hr to more than 24 hours.

Proteins may be covalently bound or noncovalently attached through nonspecific bonding. If covalent bonding between a the fusion protein and the surface is desired, the surface will usually be polyfunctional or be capable of being polyfunctionalized. Functional groups which may be present on the surface and used for linking can include carboxylic acids, aldehydes, amino groups, cyano groups, ethylenic groups, hydroxyl groups, mercapto groups and the like. The manner of linking a wide variety of compounds to various surfaces is well known and is amply illustrated in the literature.

6.2.3.1 "A Assay"Detection of PDZ-Ligand Binding Using Immobilized PL Peptide.

In one aspect, the invention provides an assay in which biotinylated candidate PL peptides are immobilized on an avidin coated surface. The binding of PDZ-domain fusion protein to this surface is then measured. In a preferred embodiment, the PDZ-domain fusion protein is a GST/PDZ fusion protein and the assay is carried out as follows:

5

(1) Avidin is bound to a surface, e.g. a protein binding surface. In one embodiment, avidin is bound to a polystyrene 96 well plate (e.g., Nunc Polysorb (cat #475094) by addition of 100 uL per well of 20 ug/mL of avidin (Pierce) in phosphate buffered saline without calcium and magnesium, pH 7.4 ("PBS", GibcoBRL) at 4°C for 12 hours. The plate is then treated to block nonspecific interactions by addition of 200 uL per well of PBS containing 2 g per 100 mL protease-free bovine serum albumin ("PBS/BSA") for 2 hours at 4°C. The plate is then washed 3 times with PBS by repeatedly adding 200 uL per well of PBS to each well of the, plate and then dumping the contents of the plate into a waste container and tapping the plate gently on a dry surface.

15

(2) Biotinylated PL peptides (or candidate PL peptides, e.g. see TABLE 4) are immobilized on the surface of wells of the plate by addition of 50 uL per well of 0.4 uM peptide in PBS/BSA for 30 minutes at 4°C. Usually, each different peptide is added to at least eight different wells so that multiple measurements (e.g. duplicates and also measurements using different (3ST/PDZ-domain fusion proteins and a GST alone negative control) can be made, and also additional negative control wells are prepared in which no peptide is immobilized. Following immobilization of the PL peptide on the surface, the plate is washed 3 times with PBS.

25

(3) GST/PDZ-domain fusion protein (prepared as described *supra*) is allowed to react with the surface by addition of 50 uL per well of a solution containing 5 ug/mL GST/PDZ-domain fusion protein in PBS/BSA for 2 hours at 4°C. As a negative control, GST alone (i.e. not a fusion protein) is added to specified wells, generally at least 2 wells (i.e. duplicate measurements) for each immobilized peptide. After the 2 hour reaction, the plate is washed 3 times with PBS to remove unbound fusion protein.

30

error in the measurement of the binding. The standard error of the mean of N measurements equals the square root of the following: the sum of the squares of the difference between each measurement and the mean, divided by the product of (N) and (N-1). Thus, in one embodiment, specific binding of the PDZ protein to the plate-bound PL peptide is determined by comparing the mean signal ("mean S") and standard error of the signal ("SE") for a particular PL-PDZ combination with the mean B1 and/or mean B2. In TABLE 2, binding was detected to be specific (denoted by an "A" in the matrix) when (1) the, mean S was at least twice the mean B1 and at least twice the mean B2 and (2) the mean S was at least six standard errors (six SE) greater than both the mean B1 and the mean B2. In addition, in the experiments summarized in TABLE 2, an additional criterion was used to ensure that none of the interactions defined as specific arose from a combined tendency of both the particular PDZ fusion protein and PL peptide tested to each give a higher than usual background. This criteria was that (3) the mean S was at least twenty times the product of the mean B 1 and the mean B2. The factor twenty times reflects that at least one of B1 and B2 is generally less than 0.1 O.D. units, and therefore twenty times the product of the mean B1 and the mean B2 is generally less than twice the mean B1 and twice the mean B2, making criteria (3) less stringent than criteria (1). Only in a few cases where the mean B1 and the mean B2 are both greater than 0.1 O.D. units (i.e. both the particular PDZ fusion protein and PL peptide tested tend to give a higher than usual background) is criteria (3) more stringent than criteria (1).

Table 4: PL Peptides			
CODE	PROTEIN NAME	GENBANK ACCESS	SEQUENCE
AA1L	Clasp-1		ISKATPALPTVSISSSAEV
AA2L	Clasp-2		ISGTPTSTMVHGMTSSSSVV
AA3L	Clasp-4		CAISGTSSDRGYGSPRYAEV
AA4L	CD3n	M33158	SVFSIPTLWSPWPPSSSSQL
AA5L-M*	CD4	M12807	SEKKTQSQPHRFQKTCSPI
AA6L	CD6	X60992	SPQPDSTDNDYDDISAA
AA7L	CD34	M81104	QATSRNGHSARQHVADTEL
AA8L	CD38	NM004334	PDKFLQCVKNPEDSSCTSEI
AA9L	CD44	M69215	QFMTADETRNLQNDMKIGV
AA10L	CD46 (Form 1)	M58050	KKGTYLTDETHREVKFTSL
AA11L	CD49E (4)	X06256	PYGTAMEKAQLKPPATSDA
AA12L	CD49F	X53586	HKAEIHAQPSDKERLTSDA
AA13L	CD95	M67454	KDITSDSENSNFRNEIQSLV
AA14L	CD97	X84700	TSGTGHNOQTRALRASESGI
AA15L	CD98	J02939	ERLKLEPHEGLLLRFPYAA
AA16L	CD105	X72012	STNHSIGSTQSTPCSTSSMA
AA17L	VCAM1	M73255	ARKANMKGYSYLVEAQSKV
AA18L	CD138	J05392	PKQANGGAYQKPTKQEEFYA
AA19L	CD148	D37781	ENLAPVTTFGKTINGYIA
AA20L	CD166	L38608	DLGNMEENKLEENNHKTEA
AA21L	CDw137 (4-1BB)	NM001561	QEEDGCSCRFPEEEEGCKEL
AA22L	DNAM-1	U56102	TREDIYVNYPTFSRRPKTRV
AA23L-M*	FasL	U11821	SSKSKSSEESQTFPGLYKL
AA25L	FceRIb	D10583	YSATYSELEDPGEMSPIDL
AA26L	Galectin3	J02921	ISKLGISGDIDLTSASYTMI
AA27L	CD114	NM000760	LNFPILQGIHVHGMALGSP
AA28L	CDW125 (IL5R)	X62156	EVICYIEKPGVETLEDVSVF
AA29.1L	CDW128A (IL8RA)	M68932	ARHRVTSYTSSSVNVSSNL
AA29.2L	CDW128B (IL8RB)	M73969	KDSRPSFVGSSSGHTSTTL
AA30L	LPAP	X81422	AWDDSARAAGGQGLHVTAL
AA31L	Mannose Receptor	NM002438	GTSDMKDLVGNIEONEHSVI
AA32L	Spectrin (beta)	NM000347	SFPFCGHRENVPGQSLVSPV
AA33L	KVI.3	AAC31761	TTNNNPNFSAVNIKIFTDV
AA34.2L	NMDA	NP000824	LNCSNRRVYKMPISIEDV
AA36L	Neuroliqin	NM018977	TFAAGFNSTGLPHSTTRV
AA37L	Glycophorin C	AAA52574	QGDPAQDAGDSRKEYFI
AA38L	Neurexin	AB011150	SSAKSSNKNKKNKDKKEYV
AA39L	Syndecan-2	A33880	GERKPSSAAYQKAPTKEFYA
AA40L	DOCK2	BAA13200	LASKSAEEGKQIPDSLSTDL
AA41L	CC CKR-1R	L09230	LERVSSSTSPSTGEHLSAGF
AA42L	CC CKR-2	U03882	GKGKSIGRAPEASLQDKEGA
AA43L	CC CKR-3	HSU28694	LERTSSVSPSTAEPELSIVF
AA44L	CC CKR-4	X85740	DTPSSSYTQSTMDHDLHDAL
AA45L	BLR-1	S56162	PSWRRSSLSESENATSLTTF
AA46L	Volt. Gated Ca2+	Q00975	SSGGRARHSYHHFPDQDHC
AA47L	CD83	Z11697	VTSFNKHLGLVTPHKTEL
AA48L	CD62E	M30640	SSSQSLES DGSYQKPSYIL
AA49L	CD5	X04391	SMQPDNSSDSYDLHGAQRL
AA55L	CD148	D37781	TIYENLAPVTTFGKTIA
AA56L	TAX	AB038239	QISPGGLEPPSEKHFRETEV
AA57L	BLR-1/CXCR5	NM001716	SWRRSSLSESENATSLTTF
AA58L	PAG	NM018440	KENDYESISDLQOGRDITRL
(PAG - Phosphoprotein Associated with GEMs)			
*The Sequence studied is mutated at positions >10 amino acids from C-terminus to increase water solubility and/or eliminate intramolecular disulfides.			

6.2.3.2 "G Assay" - Detection of PDZ-Ligand Binding Using Immobilized PDZ-Domain Fusion Polypeptide

In one aspect, the invention provides an assay in which a GST/PDZ fusion protein is immobilized on a surface ("G" assay). The binding of labeled PL peptide (as listed in **TABLE 4**) to this surface is then measured. In a preferred embodiment, the assay is carried out as follows:

(1) A PDZ-domain polypeptide is bound to a surface, e.g. a protein binding surface. In a preferred embodiment, a GST/PDZ fusion protein containing one or more PDZ domains is bound to a polystyrene 96-well plate. The GST/PDZ fusion protein can be bound to the plate by any of a variety of standard methods known to one of skill in the art, although some care must be taken that the process of binding the fusion protein to the plate does not alter the ligand-binding properties of the PDZ domain. In one embodiment, the GST/PDZ fusion protein is bound via an anti-GST antibody that is coated onto the 96-well plate. Adequate binding to the plate can be achieved when:

- a. 100 uL per well of 5 ug/mL goat anti-GST polyclonal antibody (Pierce) in PBS is added to a polystyrene 96-well plate (e.g., Nunc Polysorb) at 4°C for 12 hours.
- b. The plate is blocked by addition of 200 uL per well of PBS/BSA for 2 hours at 4°C.
- c. The plate is washed 3 times with PBS.
- d. 50 uL per well of 5 ug/mL GST/PDZ fusion protein) or, as a negative control, GST polypeptide alone (i.e. not a fusion protein) in PBS/BSA is added to the plate for 2 hours at 4°C.
- e. the plate is again washed 3 times with PBS.

(2) Biotinylated PL peptides (or candidate PL peptides, e.g. as shown in **TABLE 4**) are allowed to react with the surface by addition of 50 uL per well of 20 uM solution of the biotinylated peptide in PBS/BSA for 10 minutes at 4°C, followed by an additional 20 minute incubation at 25°C. The plate is washed 3 times with ice cold PBS.

(3) The binding of the biotinylated peptide to the GST/PDZ fusion protein surface can be detected using a variety of methods and detectors known to one of skill in the art. In one embodiment, 100 uL per well of 0.5 ug/mL streptavidin-horse radish peroxidase (HRP) conjugate dissolved in BSA/PBS is added and allowed to react for 20 minutes at 4°C.

5 The plate is then washed 5 times with 50 mM Tris pH 8.0 containing 0.2% Tween 20, and developed by addition of 100 uL per well of HRP-substrate solution (TMB, Dako) for 20 minutes at room temperature (RT). The reaction of the HRP and its substrate is terminated by addition of 100 uL per well of 1 M sulfuric acid, and the optical density (O.D.) of each well of the plate is read at 450 nm.

10

(4) Specific binding of a PL peptide and a PDZ domain polypeptide is determined by comparing the signal from the well(s) in which the PL peptide and PDZ domain polypeptide are combined, with the background signal(s). The background signal is the signal found in the negative control(s). Typically a specific or selective reaction will be at least twice
15 background signal, more typically more than 5 times background, and most typically 10 or more times the background signal. In addition, a statistically significant reaction will involve multiple measurements of the reaction with the signal and the background differing by at least two standard errors, more typically four standard errors, and most typically six or more standard errors. Correspondingly, a statistical test (e.g. a T-test) comparing repeated measurements of
20 the signal with repeated measurements of the background will result in a p-value < 0.05, more typically a p-value < 0.01, and most typically a p-value < 0.001 or less. As noted, in an embodiment of the "G" assay, the signal from binding of a given PL peptide to immobilized (surface bound) GST polypeptide alone is one suitable negative control (sometimes referred to as "B 1"). Because all measurement are done in multiples (i.e. at least duplicate) the arithmetic
25 mean (or, equivalently, average.) of several measurements is used in determining the binding, and the standard error of the mean is used in determining the probable error in the measurement of the binding. The standard error of the mean of N measurements equals the square root of the following: the sum of the squares of the difference between each measurement and the mean, divided by the product of (N) and (N-1). Thus, in one embodiment, specific binding of
30 the PDZ protein to the platebound peptide is determined by comparing the mean signal ("mean S") and standard error of the signal ("SE") for a particular PL-PDZ combination with the mean B1. In experiments summarized in **TABLE 2**, binding was determined to be specific (denoted

by a "G" in the matrix) when (1) the mean S was at least twice the mean B1 and (2) the mean S was at least six standard errors (six SE) greater than the mean B1. Results of exemplary "G" assays are shown in **Figures 1A-1D**.

6.2.3.3 "G' assay" and "G" assay"

Two specific modifications of the specific conditions described *supra* (§6.2.3.2) for the "G assay" are particularly useful. The modified assays use lesser quantities of labeled PL peptide and have slightly different biochemical requirements for detection of PDZ-ligand binding compared to the specific assay conditions described *supra*.

For convenience, the assay conditions described in this section are referred to as the "G' assay" and the "G" assay," with the specific conditions described in §6.2.3.2 being referred to as the "G⁰ assay." The "G' assay" is identical to the "G⁰ assay" except at step (2) the peptide concentration is 10 uM instead of 20 uM. This results in slightly lower sensitivity for detection of interactions with low affinity and/or rapid dissociation rate. Correspondingly, it slightly increases the certainty that detected interactions are of sufficient affinity and half-life to be of biological importance and useful therapeutic targets.

The "G" assay" is identical to the "G⁰ assay" except that at step (2) the peptide concentration is 1 uM instead of 20 uM and the incubation is performed for 60 minutes at 25°C (rather than, e.g., 10 minutes at 4°C followed by 20 minutes at 25°C). This results in lower sensitivity for interactions of low affinity, rapid dissociation rate, and/or affinity that is less at 25°C than at 4°C. Interactions will have lower affinity at 25°C than at 4°C if (as we have found to be generally true for PDZ-ligand binding) the reaction entropy is negative (i.e. the entropy of the products is less than the entropy of the reactants). In contrast, the PDZ-PL binding signal may be similar in the "G" assay" and the "G⁰ assay" for interactions of slow association and dissociation rate, as the PDZ-PL complex will accumulate during the longer incubation of the "G" assay." Thus comparison of results of the "G" assay" and the "G⁰ assay" can be used to estimate the relative entropies, enthalpies, and kinetics of different PDZ-PL interactions. (Entropies and enthalpies are related to binding affinity by the equations $\Delta G = RT \ln(K_d)$ = $\Delta H - T \Delta S$ where ΔG , H , and S are the reaction free energy, enthalpy, and entropy respectively, T is the temperature in degrees Kelvin, R is the gas constant, and K_d is the equilibrium dissociation constant). In particular, interactions that are detected only or much more strongly in the "G⁰ assay" generally have a rapid dissociation rate at 25°C ($t_{1/2} < 10$

minutes) and a negative reaction entropy, while interactions that are detected similarly strongly in the “G” assay” generally have a slower dissociation rate at 25°C ($t_{1/2} > 10$ minutes). Rough estimation of the thermodynamics and kinetics of PDZ-PL interactions (as can be achieved via comparison of results of the “G⁰ assay” versus the “G” assay” as outlined *supra*) can be used in the design of efficient inhibitors of the interactions. For example, a small molecule inhibitor based on the chemical structure of a PL that dissociates slowly from a given PDZ domain (as evidenced by similar binding in the “G” assay” as in the “G⁰ assay”) may itself dissociate slowly and thus be of high affinity.

In this manner, variation of the temperature and duration of step (2) of the “G assay” can be used to provide insight into the kinetics and thermodynamics of the PDZ-ligand binding reaction and into design of inhibitors of the reaction.

6.2.4 Assay Variations

As discussed *supra*, it will be appreciated that many of the steps in the above-described assays can be varied, for example, various substrates can be used for binding the PL and PDZ-containing proteins; different types of PDZ containing fusion proteins can be used; different labels for detecting PDZ/PL interactions can be employed; and different ways of detection can be used.

The PDZ-PL detection assays can employ a variety of surfaces to bind the PL and PDZ-containing proteins. For example, a surface can be an “assay plate” which is formed from a material (e.g. polystyrene) which optimizes adherence of either the PL protein or PDZ-containing protein thereto. Generally, the individual wells of the assay plate will have a high surface area to volume ratio and therefore a suitable shape is a flat bottom well (where the proteins of the assays are adherent). Other surfaces include, but are not limited to, polystyrene or glass beads, polystyrene or glass slides, and alike.

For example, the assay plate can be a “microtiter” plate. The term “microtiter” plate when used herein refers to a multiwell assay plate, e.g., having between about 30 to 200 individual wells, usually 96 wells. Alternatively, high density arrays can be used. Often, the individual wells of the microtiter plate will hold a maximum volume of about 250 ul. Conveniently, the assay plate is a 96 well polystyrene plate (such as that sold by Becton Dickinson Labware, Lincoln Park, N.J.), which allows for automation and high throughput screening. Other surfaces include polystyrene microtiter ELISA plates such as that sold by

Nunc Maxisorp, Inter Med, Denmark. Often, about 50 ul to 300 ul, more preferably 100 ul to 200 ul, of an aqueous sample comprising buffers suspended therein will be added to each well of the assay plate.

The detectable labels of the invention can be any detectable compound or composition which is conjugated directly or indirectly with a molecule (such as described above). The label can be detectable by itself (e.g., radioisotope labels or fluorescent labels) or, in the case of an enzymatic label, can catalyze a chemical alteration of a substrate compound or composition which is detectable. The preferred label is an enzymatic one which catalyzes a color change of a non-radioactive color reagent.

Sometimes, the label is indirectly conjugated with the antibody. One of skill is aware of various techniques for indirect conjugation. For example, the antibody can be conjugated with biotin and any of the categories of labels mentioned above can be conjugated with avidin, or vice versa (see also "A" and "G" assay above). Biotin binds selectively to avidin and thus, the label can be conjugated with the antibody in this indirect manner. See, Ausubel, *supra*, for a review of techniques involving biotin-avidin conjugation and similar assays. Alternatively, to achieve indirect conjugation of the label with the antibody, the antibody is conjugated with a small hapten (e.g. digoxin) and one of the different types of labels mentioned above is conjugated with an anti-hapten antibody (e.g. anti-digoxin antibody). Thus, indirect conjugation of the label with the antibody can be achieved.

Assay variations can include different washing steps. By "washing" is meant exposing the solid phase to an aqueous solution (usually a buffer or cell culture media) in such a way that unbound material (e.g., non-adhering cells, non-adhering capture agent, unbound ligand, receptor, receptor construct, cell lysate, or HRP antibody) is removed therefrom. To reduce background noise, it is convenient to include a detergent (e.g., Triton X) in the washing solution. Usually, the aqueous-washing solution is decanted from the wells of the assay plate following washing. Conveniently, washing can be achieved using an automated washing device. Sometimes, several washing steps (e.g., between about 1 to 10 washing steps) can be required.

Various buffers can also be used in PDZ-PL detection assays. For example, various blocking buffers can be used to reduce assay background. The term "blocking buffer" refers to an aqueous, pH buffered solution containing at least one blocking compound which is able to bind to exposed surfaces of the substrate which are not coated with a PL or PDZ-

containing protein. The blocking compound is normally a protein such as bovine serum albumin (BSA), gelatin, casein or milk powder and does not cross-react with any of the reagents in the assay. The block buffer is generally provided at a pH between about 7 to 7.5 and suitable buffering agents include phosphate and TRIS.

5 Various enzyme-substrate combinations can also be utilized in detecting PDZ-PL interactions. Examples of enzyme-substrate combinations include, for example:

(i) Horseradish peroxidase (HRPO) with hydrogen peroxide as a substrate, wherein the hydrogen peroxidase oxidizes a dye precursor (e.g. orthophenylene diamine [OPD] or 3,3',5,5'-tetramethyl benzidine hydrochloride [TMB]) (as described above).

10 (ii) alkaline phosphatase (AP) with para-Nitrophenyl phosphate as chromogenic substrate.

(iii) β -D-galactosidase (β D-Gal) with a chromogenic substrate (e.g. p-nitrophenyl- β -D-galactosidase) or fluorogenic substrate 4-methylumbelliferyl- β -D-galactosidase.

15 Numerous other enzyme-substrate combinations are available to those skilled in the art. For a general review of these, see U.S. Pat. Nos. 4,275,149 and 4,318,980, both of which are herein incorporated by reference.

Further, it will be appreciated that, although, for convenience, the present discussion primarily refers antagonists of PDZ-PL interactions, agonists of PDZ-PL interactions
20 can be identified using the methods disclosed herein or readily apparent variations thereof.

6.2.5 Results of PDL-PL Interaction Assays

TABLE 2, *supra*, shows the results of assays in which specific binding was detected using the "A" and "G" assays described herein. The top row of the table specifies the
25 source of the PDZ domain used in the GST-PDZ fusion proteins (see **TABLE 3**). The first column lists the cell surface proteins from which C-terminal peptide sequences were derived and the second column ("code") identifies the peptide used in the assay (see **TABLE 4**). The third column, "Seq" provides the sequence of the four (4) C-terminal residues of the cell surface protein and peptide. In the matrix, "A" indicates specific binding as detected in the "A" assay. "G" indicates specific binding as shown in the "G" assay. A blank indicates that no
30 specific binding was detected using the "A" or "G" assays. An asterisk (*) indicates that a pairwise interaction between the PDZ protein and the cell surface protein (or subdomains of

either) has been described by others.

6.2.5.1 New PL Motifs

As noted *supra*, TABLE 2 shows the results of assays (referred to as "PRISM MATRIX") to detect binding between PDZ proteins and candidate PL peptides. A number of specific PDZ-PL interactions are identified by the MATRIX and key amino acids and positions important in PDZ binding ("PL motifs") are deduced from these results. Not only is the MATRIX useful to catalog comprehensively PDZ-PL binding combinations, the assay can further aid in the rapid discovery and characterization of novel PL proteins and PL motifs to help in rational drug design and synthesis of PL-PDZ interaction inhibitors.

Other investigators have reported certain PL motifs important in PDZ binding, e.g., the C-terminal motifs S/T-X-V/I/L (for DLG1) and Y/F-Y/F-I/L/F for MPP1 (see, Doyle et al., 1996, *Cell* 85, 1067; Songyang et al., 1997, *Science* 275, 73). However, the reported motifs are not sufficiently specific (i.e. a large number of proteins meet these criteria yet are not necessarily actual PDZ ligands) and cover only a small number of PDZ proteins (approximately 10). The PRISM MATRIX can be used to determine ligand specificity and to deduce ligand binding motifs for any PDZ protein because it can precisely determine sequences of amino acids that do or do not result in specific PDZ binding. In addition, the assay has revealed a significant of new PDZ domain binding motifs (i.e. PL motifs): C-terminal sequence of CD6, ISAA (SEQ ID NO: 14); C- terminal sequence of CD49E, TSDA (SEQ ID NO: 24); C-terminal sequence of CD49F, TSDA (SEQ ID NO: 24); C-terminal sequence of Clasp-1, SAEV (SEQ ID NO: 175); C- terminal sequence of CLASP-4, YAEV (SEQ ID NO: 192); C-terminal sequence of CD44, KIGV (SEQ ID NO: 104); C- terminal sequence of Fas Ligand, LYKL (SEQ ID NO: ____); C-terminal sequence of IL5R, DSVF (SEQ ID NO: 94); C-terminal sequence of BLR-1, LTTF (SEQ ID NO: 217). Identification of these novel PL sequences allows the definition of novel PL motifs (See TABLE 5A, *infra*). The specificity with which these novel motifs are defined is enhanced by the fact that the MATRIX reports both positive results (i.e. PDZ-PL) combinations that result in specific binding interactions) and negative results (i.e. PDZ-PL combinations that do not result in specific binding). For example, the C-terminal sequence of CD6, SAA and the C-terminal sequence of CD49E, SDA bind to the PDZ-domain polypeptide 41.8 while the related C-terminal sequence of CD166, TEA and C- terminal sequence of CD148, YIA do not. This identifies the novel PL motif (Motif 1, *infra*)

of polypeptides terminating in alanine with serine at the -2 position and excludes polypeptides with threonine and tyrosine at the -2 position. This motif is therefore more specific than most previously identified motifs. Other novel motifs are described in **TABLE 5**.

TABLE 5

5	Position:	-3	-2	-1	C- terminal
	Motif 1	X	S	X	A
	Motif 2	X	A	D/E	V
	Motif 3	X	V/I/L	X*	V
	Motif 4	X	S/T	X	F
10	Motif 5	X	Y	X*	L

X* is any non-aromatic amino acid (any residue other than Y, F or W).

6.2.5.2 PDZ-Specific PL Motifs

15 The invention provides a method for identifying a PDZ-domain binding protein (PDZ ligand or PL) that binds to a specified PDZ protein. According to the method, a plurality of putative PL peptides (e.g., peptides or polypeptides that include at or near their carboxy terminus a sequence of the C-terminus of a naturally occurring protein known or suspected of being a PL protein, i.e., binding to at least one PDZ domain) are provided. Binding assays, typically the A and G assays as described elsewhere herein, are carried out to identify peptides that do and do not bind the specified PDZ protein (e.g., by detecting binding to a PDZ domain sequence from the specified PDZ protein). Usually, a large number of putative PL peptides are screened, as shown in **Table 2**. Thus, typically at least 2, more often at least 3, PLs that bind the specified PDZ are identified, and typically at least a plurality (e.g., in this context, at least 20 10, more often at least 20, and typically at least 40 or more) PLs that do not bind the PDZ are identified.

The sequences of the binding and nonbinding peptides are compared, for example as described *infra*, and a motif(s) characteristic of peptides that bind the PDZ domain sequence and not characteristic of peptides that do not bind the PDZ domain sequence is determined. To identify a PL protein that binds the specified PDZ protein, known proteins are examined to identify sequences (typically at or near the c-terminus, e.g., within 1, 2 or 3 residues of the c-terminus) that match the motif identified. In addition, the search parameters may include other characteristics of the protein sequences being searched, such as an expression property (e.g., expression in a particular cell type, e.g., lymphocytes, or disease state), a

functional property (e.g., receptor activity), and/or a structural property (e.g., similarity to a reference sequence). Usually this identification is carried out, at least in part, by a computer-implemented search of a database such as GenBank for proteins having the specified motif, although comparison can be made manually, particularly when the search is limited to a specific class of putative PLs.

In embodiments, the assay also includes the further step of characterizing or confirming the binding properties of the identified PL(s) and PDZ, typically by carrying out *in vivo* or *in vitro* binding assays described herein (e.g., the G assay) or known in the art (e.g., precipitation assays).

In an embodiment, a PRISM MATRIX (i.e., the representation of PL-PDZ binding interactions, e.g., interactions occurring in lymphocytes, e.g., as shown in TABLE 2) is used to identify C-terminal peptide sequence motifs characteristic of PLs (i.e., sequences that mediate binding to a particular PDZ domain-containing protein).

In an embodiment, the MATRIX is specifically arranged to facilitate identification of these motifs. To this end, the PL ligands in the MATRIX shown in TABLE 2 are ordered on the basis of C-terminal amino acid similarity, with weight given to residues reported to be important in PDZ binding (Doyle et al., 1996, *Cell* 85, 1067). In particular, in TABLE 2, peptides are first ordered based on the most C-terminal residues (zero position) in the following amino acid order: G, A, C, S, T, N, Q, D, E, H, K, R, V, I, L, M, P, F, Y, W.

Among peptides with identical C-termini, the same ranking scheme was then applied to the next most important residue for peptide binding, the -2 position, followed by the -1 position and the -3 position.

(The PDZ domains of each of the GST-PDZ fusion proteins in the MATRIX are also ordered based on amino acid sequence similarity, in this case based on multiple sequence alignment using the CLUSTAL software package. In an alternative approach, the GST-PDZ fusions can also be arranged to give additional weight in alignment to residues known in the art to be important for ligand binding.)

Based on the peptide ligands being ordered in this structure-based manner, one approach to obtaining PDZ-specific PL motifs from the MATRIX is as follows:

1. Begin by examining a single column in the MATRIX (corresponding to a single PDZ protein).
2. Determine if there exist within this column one or a few areas in which several interactions

are found, and other areas in which few or no interactions are found. For example, reading down the column of **TABLE 2** corresponding to the PDZ protein DLG1, there are many interactions in the region starting with the ligand CLASP-1 and the ending with the ligand CD34, and no interactions outside of this region.

- 5 3. If such region(s) are found, note the amino acid(s) present at the C-terminus in the region. These amino acids constitute the motif at the C-terminus. For example, for the PDZ protein DLG1, the C-terminal amino acids from ligand CLASP-1 to ligand CD34 range from V to L (including positive interactions with ligands terminating in I, the amino acid separating V and L in the amino acid order *supra*). Thus in the C-terminal motif for DLG1 is V/I/L.
- 10 4. To define a preliminary motif at the -2 position, identify a subdivision of the overall region, such that the subdivision contains several interactions and is surrounded by areas of no or few interactions. Note the amino acid(s) present at the -2 position in the subdivision. For example, in the case of DLG1, one subdivision of many interactions is from the ligand CDW128A to the ligand DOCK2. The amino acid(s) present at the -2 position in this subdivision are S and T (adjacent in the amino acid ordering), defining the - 2 position motif S/T.
- 15 5. To confirm the preliminary -2 position motif, test if it generalizes to most or all of the interactions found in the column. For example, in the case of DLG1, this process reveals that most ligands matching the C-terminal motif and the preliminary -2 position motif S/T do give detectable interactions. However, some positive interactions are also found with ligands having A, E, or Y at the -2 position. Thus a more general DLG1 -2 position motif is S/T/A/E/Y.
- 20 6. To define a preliminary motif at the -1 position, it is often helpful to identify vertically adjacent boxes which contain different results (i.e. one box contains interactions, the other does not). Frequently, these adjacent boxes will correspond to ligands that are identical at the C-terminal and -2 positions, but differ at the -1 position. If this is true, a motif is considered in which the amino acid found at the -1 position in the ligand that interacts with the PDZ protein is preferred at the -1 position of the motif, while the amino acid found at
- 25 the -1 position in the ligand giving no interactions is disfavored in the motif. This process is repeated for all available such adjacent boxes. For example, in the case of DLG1, one appropriate set of adjacent boxes is CD97 versus CD38, with the preferred -1 position
- 30

amino acid G and the disfavored E.

7. To confirm the -1 position motif, the set of preferred versus disfavored amino acids at the -1 position, identified in step 6 *supra*, is examined. If these amino acids fall into logical structural categories, e.g. the preferred amino acids are all hydrophilic while the disfavored amino acids are all hydrophobic, this confirms a -1 position motif. In the case of DLG1, preferred amino acids include R, E, and G, and disfavored include R and E. Since these sets overlap, there appears to be no -1 position motif.
8. Analogous steps to those preformed for the -1 position motif may also be preformed for the -3 position motif.

It will be apparent that the power of this analysis (or a computer implemented variation of this algorithm) increases with the size of the matrix (e.g., the number of PLs and PDZs tested). In TABLE 2, the PDZ domain-containing proteins (including MPP1, K807, DLG1, PSD95, NeDLG, 41 kd, WWP3) bind a sufficient number of ligands in the current MATRIX for the above algorithm to be practical.

For PDZs that bind a fewer number of ligands in the MATRIX, an alternative approach is useful:

1. List the set of amino acids found at the -3, -2, -1, and C-terminal positions in ligands that do bind to the PDZ. For example, in the case of AF6, the set at the C-terminal position is {V, V, V, V, I}, at the -1 position {E, F, R, Y, F}, at the -2 position {A, S, T, Y, Y}, at the -3 position {Y, V, K, E, E}.
2. Examining the set at the C-terminal position for chemical similarity. Define one or more motifs based on observed chemical similarity at this position. For example, in the case of AF6, the set {V, V, V, V, I} strongly suggests a motif V/I/L, as these are all of the mid-sized, aliphatic, hydrophobic amino acids.
3. Examine the other positions similarly. For example, for AF6, the set {E, F, R, Y, F} at the -1 position suggests a -1 position motif F/Y as these are both single ring, aromatic amino acids. A less stringent motif F/Y/D/E/K/R is also suggested, that would include all of the single ring, aromatic amino acids as well as all of the charged amino acids. However, it is unclear why these chemically dissimilar groups of amino acids—including positively charged, negatively charged, and hydrophobic amino acids--would each be preferred at this position. At the -2 position, possible motifs are S/T/Y (all hydroxylated amino acids) and

the less stringent motif A/S/T/Y. At the -3 position, possible motifs are D/E/K/R (all charged amino acids) or X (any amino acid) (as the set {Y, V, K, E, E} does not have a clear structural basis).

4. Revise the motifs at each position based on ligands that failed to bind to this PDZ. To do this, assemble a net motif using the least restrictive proposed motif at each position. Next determine how many ligands fitting this least restrictive overall motif fail to bind to this PDZ. Then see if the more restrictive proposed motif at each position would succeed in preventing these ligands from being predicted to bind. If so, the more restrictive motif is adopted. For example, for AF6 the least restrictive net motif (based on the above) is X – A/S/T/Y – F/Y/D/E/K/R – V/I/L. Fifteen different ligands in TABLE 2 (ten of which do not bind AF6) are included by this motif. Restricting the motif to F/Y at the -1 position eliminates all of the false predictions, while eliminating a minority (two of five) of the accurate predictions. As noted above, this also results in a more logical structural basis for the motif. Thus X – A/S/T/Y – F/Y – V/I/L appears to be a promising motif.
5. Check whether the promising motif includes accurately predicts all of the “strong” interactions. While every interaction in the MATRIX indicates a statistically significant interaction, not every interaction is of equal strength. (In this context, “strength” refers to the affinity/half-life of the interaction, e.g., as determined by the magnitude of the binding signal in the “G assay.”) In general, “strong” interactions result in positive interactions being detected in multiple assay formats (i.e. A, G, G’, and G’’) and particularly in the “G’ or G’’ assays” (as these use the lower ligand concentration than the G assay). For example, for AF6 the “strongest” interaction appears to be with Spectrin beta. Importantly, this ligand conforms to the promising motif *supra*. Therefore, this motif can be adopted.

PDZ domain-containing proteins in the MATRIX that bind to only one or two ligands provide a special challenge for defining PL motifs. However, in certain cases the combination of a PDZ domain binding to even a single ligand combined with the failure of that PDZ to bind related ligands allows prediction of a motif specific to that PDZ. Thus, the following process can be applied to define a PDZ-specific ligand motif based on only a single ligand that binds to that PDZ:

1. Let a preliminary guess at the C-terminal motif be X – X1 – X – X2, where X1 and X2 refer

to amino acids chemically similar or identical to the amino acid found at the -2 position and zero positions respectively of the ligand that is known to bind (X is any amino acid). For example, in the case of K545 the C-terminal sequence known to bind is that of CD105, SSMA, therefore the preliminary motif is X - A/S/T/Y - X - A/S/V.

2. Place limits on the motif, such that ligands located immediately above and below the ligand that is known to bind (but do not themselves bind) are excluded from the motif. In the case of K545, this means excluding the ligand CD49F (C-terminus TSDA) which is most easily accomplished by placing a limit on the -1 position (the zero and -2 positions are identical to SSMA, and the -3 position is very similar). A logical limit on the -1 position is to hydrophobic amino acids.
3. Check the more limited motif's ability to exclude other ligands that do not bind to the PDZ of interest. Add more restrictions as necessary, comparing the results of restrictions at different ligand positions as one proceeds. For example, in the case of K545, eight ligands that failed to bind K545 fit even the more restrictive motif. There were several possible ways to restrict the motif, e.g. by eliminating S/V at the zero position, A/Y at the -2 position, hydrophobic residues other than M at the -1 position, or residues other than A/S/T/Y at the -3 position. Of these, eliminating residues other than M at the -1 position is the only single change that is sufficient at eliminating all ligands that fail to bind K545. Therefore, one sufficient motif (i.e. that only predicts the interaction which occurs) is X - A/S/T/Y - M - A/S/V.

It is important to note that motifs created based on a single ligand that binds using the above steps will often be too broad or too narrow. For example, in the case of K545 it is unclear if G or S/V are tolerated at the zero position.

As will be apparent to one skilled in the art, the above algorithms are not mutually exclusive. Principles of each algorithm can be applied by one of experience in peptide chemistry and/or structural biology in unison to derive motifs somewhat superior to those that result from application of any one of these above algorithms in isolation. Motifs derived in this manner are provided in **TABLE 6**.

PDZ-specific PL motifs derived from the MATRIX can be applied to search genomic databases for other ligands (i.e. ligands not already in the MATRIX) that bind to a specific PDZ protein. Such searches can be performed using publicly available software such

as BLAST (available at www.ncbi.nlm.nih.gov). Knowledge of these other ligands is of practical value in several respects:

1. It permits (based on knowledge of the function of these other ligands from the literature) prediction of desirable pharmacological effects of compounds that bind to the specified PDZ domain. Thereby it allows design of compounds that have these desirable pharmacological functions.
2. It permits (based on knowledge of the function of these other ligands from the literature) prediction of undesirable pharmacological effects of compounds that bind to the specified PDZ domain. Thereby it allows prediction of side effects of pharmaceutical compounds and thus facilitates design of pharmaceutical compounds without these side effects.
3. It permits biochemical assays to be performed that directly demonstrate the binding of these other ligands to the specified PDZ. Furthermore, it allows biochemical tests to be performed that determine the ability of pharmaceuticals or candidate pharmaceuticals to inhibit (or augment) differentially the binding of different ligands to the specified PDZ. Thus, knowledge of these others ligands allows drug screens to be performed for pharmaceuticals that effect the binding of specified ones of these other ligands to the given PDZ, without effecting the binding of other ligands to the same PDZ. Thus, it allows the discovery of pharmaceuticals that inhibit only the interaction of specified PDZ-PL pairs, and thus have highly specific biological effects.
4. It permits prediction of macromolecular complexes composed of the specified PDZ domain-containing protein and ligands that bind to different domains of that protein.

Table 6 shows exemplary PL motifs derived from the PRISM MATRIX according to the invention.

TABLE 6

PL Motifs

PDZ	Motif	Comments
CASK	X-Y-X-V	Previously known
MPP1	X-S/T/Y/I-X-V	
LIMK1	X-S/T/Y-X-V	Several ligands fitting this motif do not bind LIMK1
K303	X-S-X-V	Several ligands fitting this motif do not bind K303
K807	X1-S/T-X2-V/I/L/F Preferred: X1=D/S/T; X2=D/E/N/Q/S/T	L preferred to V at C-terminus based on strength of responses (data not shown)
DLG1, PSD95, NeDLG	X-S/T/Y/A/E-X-V/I/L	Generally consistent with scientific literature; however, A/E also acceptable at -2 position
SNTa1	X-S/T/Y-D/Y-V/I/L	Both residues (D/Y) preferred at -1 position interact productively with positively charged groups
TAX- IP43	X1-S/T/Y-X-V/I Preferred: X1=E	D may be equivalent to E at -3 position; D/E may be preferred at -1 position
LDP	X-A/S-X2-V/I Preferred: X2=E	
LIM	X-S/T-X2-A/V Preferred: X2=M/R/K	Residues with a long, flexible side chain are referred at -1 position,
MINT1	X-A/S/T/I/Y-X- V/I/L/F	Many ligands give high signals with this PDZ for unclear reasons;
K545	X-A/S/T/Y-M-A/S/V	
TAX- IP2	X-S-D/E-V	
MPP2	X-S/T/Y-X-A/V/I	Several ligands fitting this motif do not bind MPP2
TIP-1	X-S/T-X2-V/I/L Possibly preferred: X2=D/E/N/Q	Several ligands fitting this motif do not bind TIP-1
PTN-4	X1-S/T-X-V/F Preferred: X1=D/E	Several ligands fitting this motif do not bind PTN-4
prIL16	D/E/K/R-V/I/L/F/Y-X- V	
CBP	X-S/T-F/Y-V	
41	X-A/S/T/Y/F-X- A/V/I/L	Small hydrophobic residue (especially V) preferred at C-terminus
AF6	X-A/S/T/Y-F/Y-V/I/L	Preferred C-terminal sequence is F-V
RGS12	X1-S/T/Y-X-V/F Preferred: X1=D/E	
PDZK1	X-T-X-F	
DLG5	X-S/T-X-V	Several ligands fitting this motif do not bind DLG5
Synt	X1-V/I/L-X2-V Possibly preferred: X1=K/R; X2=G	
WWP3	X-S/T-X2-V Preferred: X2=F/R	Both residues (F/R) preferred at -1 position have extensive hydrophobic portions
TAX- IP40	X-Y-X-V	
TIAM1	NONE	Few interactions found
DVL1	X-S/T/Y-X-V	Several ligands fitting this motif do not bind DVL1
K561	X-S/T/Y-X-V/I/L/F	Several ligands fitting this motif do not bind K561

Key: Capital letter are amino acids in standard single letter code; X refers to any amino acid; X1 or X2 refers to any amino acid, but with a preference for the amino acids stated in the table. Sequences in the motif column refer to the C-terminal four amino acids of the ligand binding to the PDZ stated in the left-most column of the table. Ligand amino acid positions (starting from the -3 position) are separated by hyphens; slashes separate various possible amino acids at a given position.

6.3 Measurement of PDZ-Ligand Binding Affinity

The “A” and “G” assays of the invention can be used to determine the “apparent affinity” of binding of a PDZ ligand peptide to a PDZ-domain polypeptide. Apparent affinity is determined based on the concentration of one molecule required to saturate the binding of a second molecule (e.g., the binding of a ligand to a receptor). Two particularly useful approaches for quantitation of apparent affinity of PDZ-ligand binding are provided *infra*.

(1) A GST/PDZ fusion protein, as well as GST alone as a negative control, are bound to a surface (e.g., a 96-well plate) and the surface blocked and washed as described supra for the “G” assay.

(2) 50 uL per well of a solution of biotinylated PL peptide (e.g. as shown in TABLE 4) is added to the surface in increasing concentrations in PBS/BSA (e.g. at 0.1 uM, 0.33 uM, 1 uM, 3.3 uM, 10 uM, 33 uM, and 100 uM). In one embodiment, the PL peptide is allowed to react with the bound GST/PDZ fusion protein (as well as the GST alone negative control) for 10 minutes at 4°C followed by 20 minutes at 25°C. The plate is washed 3 times with ice cold PBS to remove unbound labeled peptide.

(3) The binding of the PL peptide to the immobilized PDZ-domain polypeptide is detected as described supra for the “G” assay.

(4) For each concentration of peptide, the net binding signal is determined by subtracting the binding of the peptide to GST alone from the binding of the peptide to the GST/PDZ fusion protein. The net binding signal is then plotted as a function of ligand concentration and the plot is fit (e.g. by using the Kaleidagraph software package curve fitting algorithm) to the following equation, where “Signal_[ligand]” is the net binding signal at PL peptide concentration “[ligand],” “Kd” is the apparent affinity of the binding event, and “Saturation Binding” is a constant determined by the curve fitting algorithm to optimize the fit to the experimental data:

$$\text{Signal}_{[\text{ligand}]} = \text{Saturation Binding} \times ([\text{ligand}] / ([\text{ligand}] + K_d))$$

For reliable application of the above equation it is necessary that the highest peptide ligand concentration successfully tested experimentally be greater than, or at least similar to, the calculated K_d (equivalently, the maximum observed binding should be similar to the calculated saturation binding). In cases where satisfying the above criteria proves difficult, an alternative approach (infra) can be used.

The results obtained when using approach 1 are demonstrated in **FIGURES 2A and 2B**. **FIGURE 2** shows varying concentrations of biotinylated CLASP-2 (**FIG. 2A**) or Fas (**FIG. 2B**). C-terminal peptides reacted with immobilized (plate bound) GST polypeptide or GST/PDZ fusion proteins (GST/DLG1, GST/NeDLG, and GDT/PSD95) in duplicate. The signals were normalized, plotted and fit to a saturation binding curve, yielding an apparent affinity of 21 μ M for DLG1-CLASP-2 interaction, 7.5 μ M for NeDLG-CLASP-2 interaction, 45 μ M for PSD95-CLASP-2 interaction, and 54 μ M for DLG1-Fas interaction, 54 μ M for NeDLG-Fas interaction, and 85 μ M for PSD95-Fas interaction.

Approach 2:

(1) A fixed concentration of a PDZ-domain polypeptide and increasing concentrations of a labeled PL peptide (labeled with, for example, biotin or fluorescein, see TABLE 4 for representative peptide amino acid sequences) are mixed together in solution and allowed to react. In one embodiment, preferred peptide concentrations are 0.1 μ M, 1 μ M, 10 μ M, 100 μ M, 1 mM. In various embodiments, appropriate reaction times can range from 10 minutes to 2 days at temperatures ranging from 4°C to 37°C. In some embodiments, the identical reaction can also be carried out using a non-PDZ domain-containing protein as a control (e.g., if the PDZ-domain polypeptide is fusion protein, the fusion partner can be used).

(2) PDZ-ligand complexes can be separated from unbound labeled peptide using a variety of methods known in the art. For example, the complexes can be separated using high performance size-exclusion chromatography (HPSEC, gel filtration) (Rabinowitz et al., 1998, *Immunity* 9:699), affinity chromatography (e.g. using glutathione Sepharose beads), and affinity absorption (e.g., by binding to an anti-GST-coated plate as described *supra*).

(3) The PDZ-ligand complex is detected based on presence of the label on the peptide ligand using a variety of methods and detectors known to one of skill in the art. For example, if the label is fluorescein and the separation is achieved using HPSEC, an in-line fluorescence detector can be used. The binding can also be detected as described *supra* for the

G assay.

(4) The PDZ-ligand binding signal is plotted as a function of ligand concentration and the plot is fit. (e.g., by using the Kaleidagraph software package curve fitting algorithm) to the following equation, where "Signal_[ligand]" is the binding signal at PL peptide concentration "[ligand]," "K_d" is the apparent affinity of the binding event, and "Saturation Binding" is a constant determined by the curve fitting algorithm to optimize the fit to the experimental data:

$$\text{Signal}_{[\text{Ligand}]} = \text{Saturation Binding} \times ([\text{ligand}] / ([\text{ligand}] + K_d))$$

Measurement of the affinity of a labeled peptide ligand binding to a PDZ-domain polypeptide n is useful because knowledge of the affinity (or apparent affinity) of this interaction allows rational design of inhibitors of the interaction with known potency (See EXAMPLE 2). The potency of inhibitors in inhibition would be similar to (i.e. within one-order of magnitude of) the apparent affinity of the labeled peptide ligand binding to the PDZ-domain.

Thus, in one aspect, the invention provides a method of determining the apparent affinity of binding between a PDZ domain and a ligand by immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface, contacting the immobilized polypeptide with a plurality of different concentrations of the ligand, determining the amount of binding of the ligand to the immobilized polypeptide at each of the concentrations of ligand, and calculating the apparent affinity of the binding based on that data. Typically, the polypeptide comprising the PDZ domain and a non-PDZ domain is a fusion protein. In one embodiment, the e.g., fusion protein is GST-PDZ fusion protein, but other polypeptides can also be used (e.g., a fusion protein including a PDZ domain and any of a variety of epitope tags, biotinylation signals and the like) so long as the polypeptide can be immobilized In an orientation that does not abolish the ligand binding properties of the PDZ domain, e.g., by tethering the polypeptide to the surface via the non-PDZ domain via an anti-domain antibody and leaving the PDZ domain as the free end. It was discovered, for example, reacting a PDZ-GST fusion polypeptide directly to a plastic plate provided suboptimal results. The calculation of binding affinity itself can be determined using any suitable equation (e.g., as shown *supra*; also see Cantor and Schimmel (1980) BIOPHYSICAL CHEMISTRY WH Freeman & Co., San Francisco) or software.

Thus, in a preferred embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-PDZ domain (e.g., an anti-GST antibody when a GST-PDZ fusion polypeptide is used). In a preferred embodiment, the step of contacting the ligand and PDZ-domain polypeptide is carried out under the conditions provided *supra* in the description of the "G" assay. It will be appreciated that binding assays are conveniently carried out in multiwell plates (e.g., 24-well, 96-well plates, or 384 well plates).

The present method has considerable advantages over other methods for measuring binding affinities PDZ-PL affinities, which typically involve contacting varying concentrations of a GST-PDZ fusion protein to a ligand-coated surface. For example, some previously described methods for determining affinity (e.g., using immobilized ligand and GST-PDZ protein in solution) did not account for oligomerization state of the fusion proteins used, resulting in potential errors of more than an order of magnitude.

Although not sufficient for quantitative measurement of PDZ-PL binding affinity, an estimate of the relative strength of binding of different PDZ-PL pairs can be made based on the absolute magnitude of the signals observed in the "G assay." This estimate will reflect several factors, including biologically relevant aspects of the interaction, including the affinity and the dissociation rate. For comparisons of different ligands binding to a given PDZ domain-containing protein, differences in absolute binding signal likely relate primarily to the affinity and/or dissociation rate of the interactions of interest.

6.4 Assays to Identify Novel PDZ Domain Binding Moieties and to Identify Inhibitors of PDZ Protein-PL Protein Binding

Although described *supra* primarily in terms of identifying interactions between PDZ-domain polypeptides and PL proteins, the assays described *supra* and other assays can also be used to identify the binding of other molecules (e.g., peptide mimetics, small molecules, and the like) to PDZ domain sequences. For example, using the assays disclosed herein, combinatorial and other libraries of compounds can be screened, e.g., for molecules that specifically bind to PDZ domains in hematopoietic cells. Screening of libraries can be accomplished by any of a variety of commonly known methods. See, e.g., the following references, which disclose screening of peptide libraries: Parmley and Smith, 1989, *Adv. Exp. Med. Biol.* 251:215-218; Scott and Smith, 1990, *Science* 249:386-390; Fowlkes et al., 1992;

BioTechniques 13:422-427; Oldenburg et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:5393-5397; Yu et al., 1994, *Cell* 76:933-945; Staudt et al., 1988, *Science* 241:577-580; Bock et al., 1992, *Nature* 355:564-566; Tuerk et al., 1992, *Proc. Natl. Acad. Sci. USA* 89:6988-6992; Ellington et al., 1992, *Nature* 355:850-852; U.S. Patent No. 5,096,815, U.S. Patent No. 5,223,409, and
5 U.S. Patent No. 5,198,346, all to Ladner et al.; Rebar and Pabo, 1993, *Science* 263:671-673; and PCT Publication No. WO 94/18318.

In a specific embodiment, screening can be carried out by contacting the library members with a hematopoietic cell PDZ-domain polypeptide immobilized on a solid support (e.g. as described *supra* in the "G" assay) and harvesting those library members that bind to the
10 protein. Examples of such screening methods, termed "panning" techniques are described by way of example in Parmley and Smith, 1988, *Gene* 73:305-318; Fowlkes et al., 1992, *BioTechniques* 13:422-427; PCT Publication No. WO 94/18318; and in references cited hereinabove.

In another embodiment, the two-hybrid system for selecting interacting proteins
15 in yeast (Fields and Song, 1989, *Nature* 340:245-246; Chien et al., 1991, *Proc. Natl. Acad. Sci. USA* 88:9578-9582) can be used to identify molecules that specifically bind to a PDZ domain-containing protein. Furthermore, the identified molecules are further tested for their ability to inhibit transmembrane receptor interactions with a PDZ domain.

In one aspect of the invention, antagonists of an interaction between a PDZ
20 protein and a PL protein are identified. In one embodiment, a modification of the "A" assay described *supra* is used to identify antagonists. In one embodiment, a modification of the "G" assay described *supra* is used to identify antagonists.

In one embodiment, screening assays are used to detect molecules that specifically bind to PDZ domains in hematopoietic cells. Such molecules are useful as agonists
25 or antagonists of PDZ-protein-mediated cell function (e.g., cell activation, e.g., T cell activation, vesicle transport, cytokine release, growth factors, transcriptional changes, cytoskeleton rearrangement, cell movement, chemotaxis, and the like). In one embodiment, such assays are performed to screen for leukocyte activation inhibitors for drug development.

The invention thus provides assays to detect molecules that specifically bind to PDZ domain-
30 containing proteins in hematopoietic cells. For example, recombinant cells expressing PDZ domain-encoding nucleic acids can be used to produce PDZ domains in these assays and to screen for molecules that bind to the domains. Molecules are contacted with the PDZ domain

(or fragment thereof) under conditions conducive to binding, and then molecules that specifically bind to such domains are identified. Methods that can be used to carry out the foregoing are commonly known in the art.

5 It will be appreciated by the ordinarily skilled practitioner that, in one embodiment, antagonists are identified by conducting the A or G assays in the presence and absence of a known or candidate antagonist. When decreased binding is observed in the presence of a compound, that compound is identified as an antagonist. Increased binding in the presence of a compound signifies that the compound is an agonist.

10 For example, in one assay, a test compound can be identified as an inhibitor (antagonist) of binding between a PDZ protein and a PL protein by contacting a PDZ domain polypeptide and a PL peptide in the presence and absence of the test compound, under conditions in which they would (but for the presence of the test compound) form a complex, and detecting the formation of the complex in the presence and absence of the test compound. It will be appreciated that less complex formation in the presence of the test compound than
15 in the absence of the compound indicates that the test compound is an inhibitor of a PDZ protein-PL protein binding. In various embodiments, the PL peptide comprises an amino acid sequence substantially identical to the C-terminal sequence of a PL protein (e.g., CD6, CD49E, CD49F, CD138, Clasp-1, Clasp-4, VCAM1, Clasp-2, CD95, DNAM-1, CD83, CD44, CD4, CD97, Neurexin, CD3n, DOCK2, CD34, FcεRIβ, or FasLigand).

20 In one embodiment, the "G" assay is used in the presence or absence of an candidate inhibitor. In one embodiment, the "A" assay is used in the presence or absence of a candidate inhibitor.

In one embodiment (in which a G assay is used), one or more PDZ domain-containing GST-fusion proteins are bound to the surface of wells of a 96-well plate as described
25 *supra* (with appropriate controls including nonfusion GST protein). All fusion proteins are bound in multiple wells so that appropriate controls and statistical analysis can be done. A test compound in BSA/PBS (typically at multiple different concentrations) is added to wells. Immediately thereafter, 30 uL of a detectably labeled (e.g., biotinylated) peptide known to bind to the relevant PDZ domain (see, e.g., **TABLE 2**) is added in each of the wells at a final
30 concentration of, e.g., between about 2 uM and about 40 uM, typically 5 uM, 15 uM, or 25 uM. This mixture is then allowed to react with the PDZ fusion protein bound to the surface for 10 minutes at 4°C followed by 20 minutes at 25°C. The surface is washed free of unbound peptide

three times with ice cold PBS and the amount of binding of the peptide in the presence and absence of the test compound is determined. Usually, the level of binding is measured for each set of replica wells (e.g. duplicates) by subtracting the mean GST alone background from the mean of the raw measurement of peptide binding in these wells.

5 In an alternative embodiment, the A assay is carried out in the presence or absence of an test candidate to identify inhibitors of PL-PDZ interactions.

In one embodiment, a test compound is determined to be a specific inhibitor of the binding of the PDZ domain (P) and a PL (L) sequence when, at a test compound concentration of less than or equal to 1 mM (e.g., less than or equal to: 500 uM, 100 uM, 10
10 uM, 1 uM, 100 nM or 1 nM) the binding of P to L in the presence of the test compound less than about 50% of the binding in the absence of the test compound. (in various embodiments, less than about 25%, less than about 10%, or less than about 1%). Preferably, the net signal of binding of P to L in the presence of the test compound plus six (6) times the standard error of the signal in the presence of the test compound is less than the binding signal in the absence of
15 the test compound.

In one embodiment, assays for an inhibitor are carried out using a single PDZ protein-PL protein pair (e.g., a PDZ domain fusion protein and a PL peptide). In a related embodiment, the assays are carried out using a plurality of pairs, such as a plurality of different pairs listed in **TABLE 2**.

20 In some embodiments, it is desirable to identify compounds that, at a given concentration, inhibit the binding of one PL-PDZ pair, but do not inhibit (or inhibit to a lesser degree) the binding of a specified second PL-PDZ pair. These antagonists can be identified by carrying out a series of assays using a candidate inhibitor and different PL-PDZ pairs (e.g., as shown in the matrix of **TABLE 2**) and comparing the results of the assays. All such pairwise
25 combinations are contemplated by the invention (e.g., test compound inhibits binding of PL₁ to PDZ₁ to a greater degree than it inhibits binding of PL₁ to PDZ₂ or PL₂ to PDZ₂). Importantly, it will be appreciated that, based on the data provided in **TABLE 2** and disclosed herein (and additional data that can be generated using the methods described herein) inhibitors with different specificities can readily be designed.

30 For example, according to the invention, the K_i ("potency") of an inhibitor of a PDZ-PL interaction can be determined. K_i is a measure of the concentration of an inhibitor required to have a biological effect. For example, administration of an inhibitor of a PDZ-PL

interaction in an amount sufficient to result in an intracellular inhibitor concentration of at least between about 1 and about 100 Ki is expected to inhibit the biological response mediated by the target PDZ-PL interaction. In one aspect of the invention, the K_d measurement of PDZ-PL binding as determined using the methods *supra* is used in determining Ki.

5 Thus, in one aspect, the invention provides a method of determining the potency (K_i) of an inhibitor or suspected inhibitor of binding between a PDZ domain and a ligand by immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface, contacting the immobilized polypeptide with a plurality of different mixtures of the ligand and inhibitor, wherein the different mixtures comprise a fixed amount of ligand and different
10 concentrations of the inhibitor, determining the amount of ligand bound at the different concentrations of inhibitor, and calculating the K_i of the binding based on the amount of ligand bound in the presence of different concentrations of the inhibitor. In an embodiment, the polypeptide is immobilized by binding the polypeptide to an immobilized immunoglobulin that binds the non-PDZ domain. This method, which is based on the "G" assay described *supra*,
15 is particularly suited for high-throughput analysis of the K_i for inhibitors of PDZ-ligand interactions. Further, using this method, the inhibition of the PDZ-ligand interaction itself is measured, without distortion of measurements by avidity effects.

Typically, at least a portion of the ligand is detectably labeled to permit easy quantitation of ligand binding.

20 It will be appreciated that the concentration of ligand and concentrations of inhibitor are selected to allow meaningful detection of inhibition. Thus, the concentration of the ligand whose binding is to be blocked is close to or less than its binding affinity (e.g., preferably less than the 5x K_d of the interaction, more preferably less than 2x K_d, most preferably less than 1x K_d). Thus, the ligand is typically present at a concentration of less than
25 2 K_d (e.g., between about 0.01 K_d and about 2 K_d) and the concentrations of the test inhibitor typically range from 1 nM to 100 uM (e.g. a 4-fold dilution series with highest concentration 10 uM or 1 mM). In a preferred embodiment, the K_d is determined using the assay disclosed *supra*.

30 The K_i of the binding can be calculated by any of a variety of methods routinely used in the art, based on the amount of ligand bound in the presence of different concentrations of the inhibitor. in an illustrative embodiment, for example, a plot of labeled ligand binding versus inhibitor concentration is fit to the equation:

$$S_{\text{inhibitor}} = S_0 * K_i / ([I] + K_i)$$

where $S_{\text{inhibitor}}$ is the signal of labeled ligand binding to immobilized PDZ domain in the presence of inhibitor at concentration $[I]$ and S_0 is the signal in the absence of inhibitor (i.e., $[I] = 0$). Typically $[I]$ is expressed as a molar concentration.

5 In another aspect of the invention, an enhancer (sometimes referred to as, augmentor or agonist) of binding between a PDZ domain and a ligand is identified by immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface, contacting the immobilized polypeptide with the ligand in the presence of a test agent and determining the amount of ligand bound, and comparing the amount of ligand bound in the
10 presence of the test agent with the amount of ligand bound by the polypeptide in the absence of the test agent. At least two-fold (often at least 5-fold) greater binding in the presence of the test agent compared to the absence of the test agent indicates that the test agent is an agent that enhances the binding of the PDZ domain to the ligand. As noted *supra*, agents that enhance PDZ-ligand interactions are useful for disruption (dysregulation) of biological events requiring
15 normal PDZ-ligand function (e.g., cancer cell division and metastasis, and activation and migration of immune cells).

The invention also provides methods for determining the "potency" or " K_{enhancer} " of an enhancer of a PDZ- ligand interaction. For example, according to the invention, the K_{enhancer} of an enhancer of a PDZ-PL interaction can be determined, e.g., using the K_d of PDZ-
20 PL binding as determined using the methods described *supra*. K_{enhancer} is a measure of the concentration of an enhancer expected to have a biological effect. For example, administration of an enhancer of a PDZ-PL interaction in an amount sufficient to result in an intracellular inhibitor concentration of at least between about 0.1 and about 100 K_{enhancer} (e.g., between about 0.5 and about 50 K_{enhancer}) is expected to disrupt the biological response mediated by the target
25 PDZ-PL interaction.

Thus, in one aspect the invention provides a method of determining the potency (K_{enhancer}) of an enhancer or suspected enhancer of binding between a PDZ domain and a ligand by immobilizing a polypeptide comprising the PDZ domain and a non-PDZ domain on a surface, contacting the immobilized polypeptide with a plurality of different mixtures of the
30 ligand and enhancer, wherein the different mixtures comprise a fixed amount of ligand, at least a portion of which is detectably labeled, and different concentrations of the enhancer, determining the amount of ligand bound at the different concentrations of enhancer, and

calculating the potency (K_{enhancer}) of the enhancer from the binding based on the amount of ligand bound in the presence of different concentrations of the enhancer. Typically, at least a portion of the ligand is detectably labeled to permit easy quantitation of ligand binding. This method, which is based on the “G” assay described *supra*, is particularly suited for high-throughput analysis of the K_{enhancer} for enhancers of PDZ-ligand interactions.

It will be appreciated that the concentration of ligand and concentrations of enhancer are selected to allow meaningful detection of enhanced binding. Thus, the ligand is typically present at a concentration of between about 0.01 Kd and about 0.5 Kd and the concentrations of the test agent/enhancer typically range from 1 nM to 1 mM (e.g. a 4-fold dilution series with highest concentration 10 uM or 1 mM). In a preferred embodiment, the Kd is determined using the assay disclosed *supra*.

The potency of the binding can be determined by a variety of standard methods based on the amount of ligand bound in the presence of different concentrations of the enhancer or augmentor. For example, a plot of labeled ligand binding versus enhancer concentration can be fit to the equation:

$$S([E]) = S(0) + (S(0) \cdot (D_{\text{enhancer}} - 1) \cdot [E]) / ([E] + K_{\text{enhancer}})$$

where “ K_{enhancer} ” is the potency of the augmenting compound, and “ D_{enhancer} ” is the fold-increase in binding of the labeled ligand obtained with addition of saturating amounts of the enhancing compound, $[E]$ is the concentration of the enhancer. It will be understood that saturating amounts are the amount of enhancer such that further addition does not significantly increase the binding signal. Knowledge of “ K_{enhancer} ” is useful because it describes a concentration of the augmenting compound in a target cell that will result in a biological effect due to dysregulation of the PDZ-PL interaction. Typical therapeutic concentrations are between about 0.1 and about 100 K_{enhancer} .

6.4.1 Global Analysis of PDZ-PL Interactions

As described *supra*, the present invention provides powerful methods for analysis of PDZ-ligand interactions, including high-throughput methods such as the “G” assay and affinity assays described *supra*. In one embodiment of the invention, the affinity is determined for a particular ligand and a plurality of PDZ proteins. Typically the plurality is at least 5, and often at least 25, or at least 40 different PDZ proteins. In a preferred embodiment, the plurality of different PDZ proteins are from a particular tissue (e.g., central nervous system,

spleen, cardiac muscle, kidney) or a particular class or type of cell, (e.g., a hematopoietic cell, a lymphocyte, a neuron) and the like. In a most preferred embodiment, the plurality of different PDZ proteins represents a substantial fraction (e.g., typically a majority, more often at least 80%) of all of the PDZ proteins known to be, or suspected of being, expressed in the tissue or cell(s), e.g., all of the PDZ proteins known to be present in lymphocytes. In an embodiment, the plurality is at least 50%, usually at least 80%, at least 90% or all of the PDZ proteins disclosed herein as being expressed in hematopoietic cells (see Table 7).

In one embodiment of the invention, the binding of a ligand to the plurality of PDZ proteins is determined. Using this method, it is possible to identify a particular PDZ domain bound with particular specificity by the ligand. The binding may be designated as “specific” if the affinity of the ligand to the particular PDZ domain is at least 2-fold that of the binding to other PDZ domains in the plurality (e.g., present in that cell type). The binding is deemed “very specific” if the affinity is at least 10-fold higher than to any other PDZ in the plurality or, alternatively, at least 10-fold higher than to at least 90%, more often 95% of the other PDZs in a defined plurality. Similarly, the binding is deemed “exceedingly specific” if it is at least 100-fold higher. For example, a ligand could bind to 2 different PDZs with an affinity of 1 μM and to no other PDZs out of a set 40 with an affinity of less than 100 μM . This would constitute specific binding to those 2 PDZs. Similar measures of specificity are used to describe binding of a PDZ to a plurality of PLs.

It will be recognized that high specificity PDZ-PL interactions represent potentially more valuable targets for achieving a desired biological effect. The ability of an inhibitor or enhancer to act with high specificity is often desirable. In particular, the most specific PDZ-ligand interactions are also the best therapeutic targets, allowing specific inhibition of the interaction.

Thus, in one embodiment, the invention provides a method of identifying a high specificity interaction between a particular PDZ domain and a ligand known or suspected of binding at least one PDZ domain, by providing a plurality of different immobilized polypeptides, each of said polypeptides comprising a PDZ domain and a non-PDZ domain; determining the affinity of the ligand for each of said polypeptides, and comparing the affinity of binding of the ligand to each of said polypeptides, wherein an interaction between the ligand and a particular PDZ domain is deemed to have high specificity when the ligand binds an

immobilized polypeptide comprising the particular PDZ domain with at least 2-fold higher affinity than to immobilized polypeptides not comprising the particular PDZ domain.

In a related aspect, the affinity of binding of a specific PDZ domain to a plurality of ligands (or suspected ligands) is determined. For example, in one embodiment, the invention provides a method of identifying a high specificity interaction between a PDZ domain and a particular ligand known or suspected of binding at least one PDZ domain, by providing an immobilized polypeptide comprising the PDZ domain and a non-PDZ domain; determining the affinity of each of a plurality of ligands for the polypeptide, and comparing the affinity of binding of each of the ligands to the polypeptide, wherein an interaction between a particular ligand and the PDZ domain is deemed to have high specificity when the ligand binds an immobilized polypeptide comprising the PDZ domain with at least 2-fold higher affinity than other ligands tested. Thus, the binding may be designated as "specific" if the affinity of the PDZ to the particular PL is at least 2-fold that of the binding to other PLs in the plurality (e.g., present in that cell type). The binding is deemed "very specific" if the affinity is at least 10-fold higher than to any other PL in the plurality or, alternatively, at least 10-fold higher than to at least 90%, more often 95% of the other PLs in a defined plurality. Similarly, the binding is deemed "exceedingly specific" if it is at least 100-fold higher. Typically the plurality is at least 5 different ligands, more often at least 10. In an embodiment, the plurality of ligands comprises at least 1, typically at least 2, more often at least 5, and sometimes at least 10 ligands selected from CD105, VCAM1, CD95, Spectrin beta, KV1.3, DNAM1, Neuroligin 3, TAX, CD44 (long form), CD38, CD3n, LPAP, CD46 (form 1), CDw128B (IL-8 receptor B), DOCK2, PAG, CD34, BLR-1 (or a polypeptide comprising a C-terminal sequence (e.g., at least about 3, 4, 6, 8 or 10 residues) from such a ligand).

TABLE 7

PDZ Domain-Containing Genes Expressed in T Cells and B Cells

<i>PDZ gene name</i>	<i>Expressed in T / B cells</i>	<i>Genebank acc. #</i>
AF6	T- / B-cells	430993
BAI I associated prot.	T- / B-cells	3370997
CASK (mouse)	T- / B-cells	3087815
Connector enhancer	B-cells	3930780
Cytohesin bind. Prot.	T- / B-cells	3192908

DLG1	T- / B-cells	475816
DLG5 (pdlg)	T- / B-cells	3650451
DVL1	T- / B-cells	2291005
DVL3	T- / B-cells	6806886
GTPase	T- / B-cells	3004860
Guanin-exchange factor 1	T- / B-cells	6650765
hypoth. 41.8 kd	T- / B-cells	3882222
PDZ domain containing prot.	T cells only	2370148
KIAA147	T- / B-cells	1469875
KIAA0300	T- / B-cells	2224540
KIAA0303	T- / B-cells	2224546
KIAA0316	T-cells	6683123
KIAA0380	T- / B-cells	2224700
KIAA0440	T- / B-cells	2662160
KIAA0545	T- / B-cells	303617
KIAA0561	T- / B-cells	3043645
KIAA0559	B-cells	3043641
KIAA0807	T- / B-cells	3882334
KIAA0858	T- / B-cells	42402004
KIAA0902	T- / B-cells	4240304
LIMK1	T- / B-cells	4587498
LIMK2	T- / B-cells	1805593
LIM domain prot	T- / B-cells	2957144
LIM protein	T- / B-cells	3108092
MINT1	T- / B-cells	2625024
MINT3	T- / B-cells	3169808
MPP1	T- / B-cells	189785
MPP2	T- / B-cells	939884
NE-DLG	T- / B-cells	1515354
NOS1	T- / B-cells	642525
novel serine protease	T- / B-cells	1621243
PDZK1	T- / B-cells	2944188

PICK8	T- / B-cells	4678411
PTN-3	T- / B-cells	179912
PTN-4	B cells	190747
prIL16	T- / B-cells	1478492
PSD95	T- / B-cells	3318652
RPIP8	T- / B-cells	5730014
RGS12	T- / B-cells	3290015
serine protease	T- / B-cells	2738914
26s subunit p27	T-cells	9184389
hSYNTENIN	T- / B-cells	2795862
SYNTR. 1 alpha	T- / B-cells	1145727
TAX1-IP	T- / B-cells	2613001
TAX2-IP	T- / B-cells	2613003
TAX2-like protein	T- / B-cells	3253116
TAX33-IP	T- / B-cells	2613007
TAX40-IP (PAR-6)	T- / B-cells	2613011
Tax43-IP (SYN. Beta1)	T- / B-cells	2613011
TIAM	T- / B-cells	4507500
wwp3	T- / B-cells	2695619
X11 prot. beta	T- / B-cells	3005559
ZO1	T- / B-cells	292937

6.4.2 Use of Array for Global Predictions

One discovery of the present inventors relates to the important and extensive roles played by interactions between PDZ proteins and PL proteins, particularly in the biological function of hematopoietic cells and other cells involved in the immune response. Further, it has been discovered that valuable information can be ascertained by analysis (e.g., simultaneous analysis) of a large number of PDZ-PL interactions. In a preferred embodiment, the analysis encompasses all of the PDZ proteins expressed in a particular tissue (e.g., spleen) or type or class of cell (e.g., hematopoietic cell, neuron, lymphocyte, B cell, T cell and the like). Alternatively, the analysis encompasses at least about 5, or at least about 10, or at least about 12, or at least about 15 and often at least 50 different polypeptides, up to about 60, about 80,

about 100, about 150, about 200, or even more different polypeptides; or a substantial fraction (e.g., typically a majority, more often at least 80%) of all of the PDZ proteins known to be, or suspected of being, expressed in the tissue or cell(s), e.g., all of the PDZ proteins known to be present in lymphocytes. In an embodiment, the plurality is at least 50%, usually at least 80%, at least 90% or all of the PDZ proteins disclosed herein as being expressed in hematopoietic cells (see Table 7).

It will be recognized that the arrays and methods of the invention are directed to analysis of PDZ and PL interactions, and involve selection of such proteins for analysis. While the devices and methods of the invention may include or involve a small number of control polypeptides, they typically do not include significant numbers of proteins or fusion proteins that do not include either PDZ or PL domains (e.g., typically, at least about 90% of the arrayed or immobilized polypeptides in a method or device of the invention is a PDZ or PL sequence protein, more often at least about 95%, or at least about 99%).

In an embodiment the array includes at least one, preferably at least 1, more often at least 5 or at least 10 and sometimes all of the following PDZ proteins present in lymphocytes: BAI I associated prot., Connector enhancer, DLG5 (pdlg), DVL3, GTPase, Guanine-exchange factor 1, PDZ domain containing prot., KIAA147, KIAA0300, KIAA0380, KIAA0440, KIAA0545, KIAA0807, KIAA0858, KIAA0902, novel serine protease, PDZK1, PICK8, PTN-3, RPIP8, serine protease, 26s subunit p27, hSYNTENIN, TAX1-IP, TAX2-like protein, wwp3, X11 prot. beta, ZO1.

It will be apparent from this disclosure that analysis of the relatively large number of different interactions preferably takes place simultaneously. In this context, "simultaneously" means that the analysis of several different PDZ-PL interactions (or the effect of a test agent on such interactions) is assessed at the same time. Typically the analysis is carried out in a highthroughput (e.g., robotic) fashion. One advantage of this method of simultaneous analysis is that it permits rigorous comparison of multiple different PDZ-PL interactions. For example, as explained in detail elsewhere herein, simultaneous analysis (and use of the arrays described *infra*) facilitates, for example, the direct comparison of the effect of an agent (e.g., an potential interaction inhibitor) on the interactions between a substantial portion of PDZs and/or PLs in a tissue or cell.

Accordingly, in one aspect, the invention provides an array of immobilized polypeptide comprising the PDZ domain and a non-PDZ domain on a surface. Typically, the



array comprises at least about 5, or at least about 10, or at least about 12, or at least about 15 and often at least 50 different polypeptides. In one preferred embodiment, the different PDZ proteins are from a particular tissue (e.g., central nervous system, spleen, cardiac muscle, kidney) or a particular class or type of cell, (e.g., a hematopoietic cell, a lymphocyte, a neuron) and the like. In a most preferred embodiment, the plurality of different PDZ proteins represents a substantial fraction (e.g., typically a majority, more often at least 80%) of all of the PDZ proteins known to be, or suspected of being, expressed in the tissue or cell(s), e.g., all of the PDZ proteins known to be present in lymphocytes. In an embodiment, the plurality is at least 50%, usually at least 80%, at least 90% or all of the PDZ proteins disclosed herein as being expressed in hematopoietic cells (see Table 7) e.g.; all of the PDZ proteins known to be present in lymphocytes. In an embodiment, the plurality is at least 50%, usually at least 80%, at least 90% or all of the PDZ proteins disclosed herein as being expressed in hematopoietic cells (see Table 7).

In an embodiment the array includes at least one, preferably at least 1, typically at least 5 and sometimes all of the following PDZ proteins present in lymphocytes: BAI 1 associated prot., Connector enhancer, DLG5 (pdlg), DVL3, GTPase, Guanine-exchange factor 1, PDZ domain containing prot., KIAA147, KIAA0300, KIAA0380, KIAA0440, KIAA0545, KIAA0807, KIAA0858, KIAA0902, novel serine protease, PDZK1, PICK8, PTN-3, RPIP8, serine protease, 26s subunit p27, hSYNTENIN, TAX1-IP, TAX2-like protein, wwp3, X11 prot. beta, ZO1. In this context, “array” refers to an ordered series of of immobilized polypeptides in which the identity of each polypeptide is associated with its location. In some embodiments the plurality of polypeptides are arrayed in a “common” area such that they can be simultaneously exposed to a solution (e.g., containing a ligand or test agent). For example, the plurality of polypeptides can be on a slide, plate or similar surface, which may be plastic, glass, metal, silica, beads or other surface to which proteins can be immobilized. In a different embodiment, the different immobilized polypeptides are situated in separate areas, such as different wells of multi-well plate (e.g., a 24-well plate, a 96-well plate, a 384 well plate, and the like). It will be recognized that a similar advantage can be obtained by using multiple arrays in tandem.

6.4.3 Analysis of PDZ-PL Inhibition Profile

In one aspect, the invention provides a method for determining if a test compound inhibits any PDZ-ligand interaction in large set of PDZ-ligand interaction (e.g., some or all of the PDZ-ligands interactions described in Table 2; a majority of the PDZ-ligands identified in a particular cell or tissue as described *supra* (e.g., lymphocytes) and the like. In one embodiment, the PDZ domains of interest are expressed as GST-PDZ fusion proteins and immobilized as described herein. For each PDZ domain, a labeled ligand that binds to the domain with a known affinity is identified as described herein.

As disclosed herein, numerous PDZ-PL interactions occur in cells of the hematopoietic system. For any known or suspected modulator (e.g., inhibitor) of a PDL-PL interaction(s), it is useful to know which interactions are inhibited (or augmented). For example, an agent that inhibits *all* PDZ-PL interactions in a cell (e.g., a lymphocyte) will have different uses than an agent that inhibits only one, or a small number, of specific PDZ-PL interactions. The profile of PDZ interactions inhibited by a particular agent is referred to as the "inhibition profile" for the agent, and is described in detail below. The profile of PDZ interactions enhanced by a particular agent is referred to as the "enhancement profile" for the agent. It will be readily apparent to one of skill guided by the description of the inhibition profile how to determine the enhancement profile for an agent. The present invention provides methods for determining the PDZ interaction (inhibition/enhancement) profile of an agent in a single assay.

In one aspect, the invention provides a method for determining the PDZ-PL inhibition profile of a compound by providing (i) a plurality of different immobilized polypeptides, each of said polypeptides comprising a PDZ domain and a non-PDZ domain and (ii) a plurality of corresponding ligands, wherein each ligand binds at least one PDZ domain in (i), then contacting each of said immobilized polypeptides in (i) with a corresponding ligand in (ii) in the presence and absence of a test compound, and determining for each polypeptide-ligand pair whether the test compound inhibits binding between the immobilized polypeptide and the corresponding ligand.

Typically the plurality is at least 5, and often at least 25, or at least 40 different PDZ proteins. In a preferred embodiment, the plurality of different ligands and the plurality of different PDZ proteins are from the same tissue or a particular class or type of cell, e.g., a hematopoietic cell, a lymphocyte, a neuron and the like. In a most preferred embodiment, the plurality of different PDZs represents a substantial fraction (e.g., at least 80%) of all of the

PDZs known to be, or suspected of being, expressed in the tissue or cell(s), e.g., all of the PDZs known to be present in lymphocytes (for example, at least 80%, at least 90% or all of the PDZs disclosed herein as being expressed in hematopoietic cells).

In one embodiment, the inhibition profile is determined as follows: A plurality (e.g., all known) PDZ domains expressed in a cell (e.g., lymphocytes) are expressed as GST-fusion proteins and immobilized without altering their ligand binding properties as described *supra*. For each PDZ domain, a labeled ligand that binds to this domain with a known affinity is identified. If the set of PDZ domains expressed in lymphocytes is denoted by $\{P1 \dots Pn\}$, any given PDZ domain P_i binds a (labeled) ligand L_i with affinity $K_{d,i}$. To determine the inhibition profile for a test agent "compound X" the "G" assay (*supra*) can be performed as follows in 96-well plates with rows A-H and columns 1-12. Column 1 is coated with P_1 and washed. The corresponding ligand L_1 is added to each washed coated well of column 1 at a concentration $0.5 K_{d,1}$ with (rows B, D, F, H) or without (rows A, C, E, G) between about 1 and about 1000 μM of test compound X. Column 2 is coated with P_2 , and L_2 (at a concentration $0.5 K_{d,2}$) is added with or without inhibitor X. Additional PDZ domains and ligands are similarly tested.

Compound X is considered to inhibit the binding of L_i to P_i if the average signal in the wells of column i containing X is less than half the signal in the equivalent wells of the column lacking X. Thus, in this single assay one determines the full set of lymphocyte PDZs that are inhibited by compound X.

In some embodiments, the test compound X is a mixture of compounds, such as the product of a combinatorial chemistry synthesis as described *supra*. In some embodiments, the test compound is known to have a desired biological effect, and the assay is used to determine the mechanism of action (i.e., if the biological effect is due to modulating a PDZ-PL interaction).

It will be apparent that an agent that modulates only one, or a few PDZ-PL interactions, in a panel (e.g., a panel of all known PDZs lymphocytes, a panel of at least 10, at least 20 or at least 50 PDZ domains) is a more specific modulator than an agent that modulate many or most interactions. Typically, an agent that modulates less than 20% of PDZ domains in a panel (e.g., Table 2) is deemed a "specific" inhibitor, less than 6% a "very specific" inhibitor, and a single PDZ domain a "maximally specific" inhibitor.

It will also be appreciated that "compound X" may be a composition containing mixture of compounds (e.g., generated using combinatorial chemistry methods) rather than a

single compound.

Several variations of this assay are contemplated:

In some alternative embodiments, the assay above is performed using varying concentrations of the test compound X, rather than fixed concentration. This allows
5 determination of the K_i of the X for each PDZ as described above.

In an alternative embodiment, instead of pairing each PDZ P_i with a specific labeled ligand L_i , a mixture of different labeled ligands is created such that for every PDZ at least one of the ligands in the mixture binds to this PDZ sufficiently to detect the binding in the "G" assay. This mixture is then used for every PDZ domain.

10 In one embodiment, compound X is known to have a desired biological effect, but the chemical mechanism by which it has that effect is unknown. The assays of the invention can then be used to determine if compound X has its effect by binding to a PDZ domain.

In one embodiment, PDZ-domain containing proteins are classified into groups based on their biological function, e.g. into those that regulate chemotaxis versus those that
15 regulate transcription. An optimal inhibitor of a particular function (e.g., including but not limited to an anti-chemotactic agent, an anti-T cell activation agent, cell-cycle control, vesicle transport, apoptosis, etc.) will inhibit multiple PDZ-ligand interactions involved in the function (e.g., chemotaxis, activation) but few other interactions. Thus, the assay is used in one embodiment in screening and design of a drug that specifically blocks a particular function.

20 For example, an agent designed to block chemotaxis might be identified because, at a given concentration, the agent inhibits 2 or more PDZs involved in chemotaxis but fewer than 3 other PDZs, or that inhibits PDZs involved in chemotaxis with a $K_i > 10$ -fold better than for other PDZs. Thus, the invention provides a method for identifying an agent that inhibits a first selected PDZ-PL interaction or plurality of interactions but does not inhibit a second selected
25 PDZ-PL interaction or plurality of interactions. The two (or more) sets of interactions can be selected on the basis of the known biological function of the PDZ proteins, the tissue specificity of the PDZ proteins, or any other criteria. Moreover, the assay can be used to determine effective doses (i.e., drug concentrations) that result in desired biological effects while avoiding undesirable effects.

30 6.4.4 Side Effects of PDZ-PL Modulator Interactions

In a related embodiment, the invention provides a method for determining likely

side effects of a therapeutic that inhibits PDZ-ligand interactions. The method entails identifying those target tissues, organs or cell types that express PDZ proteins and ligands that are disrupted by a specified inhibitor. If, at a therapeutic dosage, a drug intended to have an effect in one organ system (e.g., hematopoietic system) disrupts PDZ-PL interactions in a different system (e.g., CNS) it can be predicted that the drug will have effects ("side effects") on the second system. It will be apparent that the information obtained from this assay will be useful in the rational design and selection of drugs that do not have the side-effect.

In one embodiment, for example, a comprehensive PDZ protein set is obtained. A "perfectly comprehensive" PDZ protein set is defined as the set of all PDZ proteins expressed in the subject animal (e.g., humans). A comprehensive set may be obtained by analysis of, for example, the human genome sequence. However, a "perfectly comprehensive" set is not required and any reasonably large set of PDZ domain proteins (e.g., the set of all known PDZ proteins; or the set listed in Table 7) will provide valuable information.

In one embodiment, the method involves some of all of the following steps:

a) For each PDZ protein, determine the tissues in which it is highly expressed. This can be done experimentally although the information generally will be available in the scientific literature;

b) For each PDZ protein (or as many as possible), identify the cognate PL(s) bound by the PDZ protein;

c) Determine the K_i at which the test agent inhibits each PDZ-PL interaction, using the methods described *supra*;

d) From this information it is possible to calculate the pattern of PDZ-PL interactions disrupted at various concentrations of the test agent

By correlating the set of PDZ-PL interactions disrupted with the expression pattern of the members of that set, it will be possible to identify the tissues likely affected by the agent.

Additional steps can also be carried out, including determining whether a specified tissue or cell type is exposed to an agent following a particular route of administration. This can be determined using basis pharmacokinetic methods and principles.

6.4.5 Modulation of Activities

The PDZ binding moieties and PDZ protein -PL protein binding antagonists of the invention are used to modulate biological activities or functions of cells (e.g., hematopoietic

cells, such as T cells and B cells and the like), endothelial cells, and other immune system cells, as described herein, and for treatment of diseases and conditions in human and nonhuman animals (e.g., experimental models). Exemplary biological activities are listed *supra*.

When administered to patients, the compounds of the invention (e.g., PL-PDZ interaction inhibitors) are useful for treating (ameliorating symptoms of) a variety of diseases and conditions, including diseases characterized by inflammatory and humoral immune responses, e.g., inflammation, allergy (e.g., systemic anaphylaxis, hypersensitivity responses, drug allergies, insect sting allergies; inflammatory bowel diseases, ulcerative colitis, ileitis and enteritis; psoriasis and inflammatory dermatoses, scleroderma; respiratory allergic diseases such as asthma, allergic rhinitis, hypersensitivity lung diseases, and the like vasculitis, rh incompatibility, transfusion reactions, drug sensitivities, PIH, atopic dermatitis, eczema, rhinitis; autoimmune diseases, such as arthritis (rheumatoid and psoriatic), multiple sclerosis, systemic lupus erythematosus, insulin-dependent diabetes, glomerulonephritis, scleroderma, MCTD, IDDM, Hashimoto thyroiditis, Goodpasture syndrome, psoriasis and the like, osteoarthritis, polyarthritis, graft rejection (e.g., allograft rejection, e.g., renal allograft rejection, graft-vs-host disease, transplantation rejection (cardiac, kidney, lung, liver, small bowel, cornea, pancreas, cadaver, autologous, bone marrow, xenotransplantation)), atherosclerosis, angiogenesis-dependent disorders, cancers (e.g., melanomas and breast cancer, prostate cancer, leukemias, lymphomas, metastatic disease), infectious diseases (e.g., viral infection, such as HIV, measles, parainfluenza, virus-mediated cell fusion,), ischemia (e.g., post-myocardial infarction complications, joint injury, kidney, scleroderma).

The PL proteins and PDZ proteins listed in TABLE 2 are well characterized, and one of skill, guided by this disclosure (including the discovery of the interactions between PL proteins and PDZ proteins described herein), will recognize many uses for modulators (e.g., enhancers or inhibitors) of PDZ-PL interactions such as those described in TABLE 2. To further assist the reader, a discussion of the characteristics of selected PL proteins (and their function) is provided *infra*. It will be recognized that this discussion is not comprehensive and is not intended to limit the invention in any way. Moreover, nothing in this section should be construed as an intention by the inventors to be limited to a particular mechanism of action.

A. CD6

As shown *supra*, CD6 binds PDZ protein 41.8. CD6 is expressed on

thymocytes, T cells, and B cell chronic lymphocytic leukemias. CD6 plays a role in T cell co-stimulation and CD6 negative T cells are less autoreactive than CD6 positive T cells. Inhibition of CD6 and CD6/41.8 interactions is predicted to reduce the symptoms of graft-versus-host disease (GVHD) or psoriasis. Thus, in one embodiment of the invention, GVHD is reduced in a patient receiving donor bone marrow cells by pre-treating the cells with an effective amount of an antagonist. In combination with post-transplantation immunosuppressive therapy such as FK506, Cellcept, or cyclosporin, CD6-PDZ interaction inhibitors will improve overall survival of transplantation patients (e.g., leukemia patients).

10 **B. CD49e (ALPHA-4)**

As shown by the experiments reported herein, the C-terminal end of CD49e binds to the PDZ-domain-containing protein 41.8 kD. CD49e is a 110 kD transmembrane membrane protein of the integrin alpha family (integrin alpha 5). Paired with the integrin beta-1 subunit it forms VLA-5. VLA-5 is expressed predominantly on hematopoietic and lymphoid lineage cells including monocytes, basophils, T cells, and activated B cells. VLA-5 is the receptor for the ubiquitously-expressed adhesion molecule fibronectin. Tissue injury such as myocardial infarction releases soluble fragments of fibronectin. Binding of these soluble fragments to VLA-5 results in chemotaxis of immune cells including monocytes to the source of fibronectin, as well as down-modulation of VLA-5 expression on these cells. Such ligand-induced down-modulation is a common and required feature of chemotactic receptors. Once immune cells migrate fully to the source of fibronectin, adhesion to the fibronectin surface is enhanced by fibronectin-VLA-5 interaction. Without intending to be bound by a particular mechanism, the 41.8/CD49e interaction is believed to be necessary for proper membrane distribution of CD49e and/or recycling of CD49e such that when it is disrupted, the migration and adherence to fibronectin-containing surfaces is similarly disrupted, resulting in an inability of immune system cells to effectively migrate toward a fibronectin source and adhere to fibronectin-containing surfaces. Such disruption would therefore result in desirable reduced inflammatory processes, including reduced post-myocardial infarction inflammation. Other diseases to be treated include but are not limited to joint inflammation, psoriasis, contact allergy, Crohn's Disease, inflammatory bowel disease, eczema, atopic dermatitis.

C. CD49F (VLA-6 α subunit)

As shown *supra*, CD49F binds PDZ protein 41.8. CD49f is known as an integrin subunit that pairs either with the $\beta 1$ integrin subunit (CD29), forming VLA-6, or with CD104 ($\beta 4$ integrin subunit). The integrin supergene family consists of a number of cell surface $\alpha\beta$ heterodimers important for many different physiologic processes, including
5 embryogenesis, thrombosis, wound healing, tumorigenesis and immune responses. Each β chain can pair with various α chains. Both VLA-6 and CD49f/CD104 are widely expressed on epithelia in non-lymphoid tissues. VLA-6 is also expressed on platelets, monocytes, thymocytes and T lymphocytes, with an increased expression on activated and resting memory T cells.

10 Inhibition of interactions between VLA-6 and 41.8 has a number of therapeutic functions such as the prevention and treatment of metastatic cancers, and treatment of overactive immunity. For example, VLA-6 is associated with invasion of prostate carcinoma and plays a role in the metastasis of breast cancer. Blockage of VLA-6 function combined with conventional treatment for prostate cancer, would be a more effective treatment by preventing
15 metastatic disease (see, Cress et al., 1995, Cancer Metastasis R). Blockage of CD49f through PDZ interaction may also treat Rh incompatibility by blunting memory response or in the treatment of keloids.

D. CD138 (syndecan-1)

20 CD138 is a transmembrane proteoglycan receptor with the extracellular domain functioning as a ligand binding domain for various extracellular matrix components and the intracellular portion functioning to alter cytoskeleton and transduce intracellular signals. CD138 also binds FGF2 and may be a co-receptor for FGF receptor (Yayon et al., 1991).

As shown *supra*, CD138 interacts with 41.8kD protein and TIAM1. The c-
25 terminus of CD138 has also been reported to bind the PDZ domains of syntenin and human CASK (Cohen et al., 1998, *J. Cell. Biol.* 142:129-138; Grootjans et al., 1997, *PNAS* 94:13683-13688; Hsueh et al., 1998, *J. Cell. Biol.* 142:139-151). CD138 is expressed in pre-B cells, immature B cells, plasma cells, neural cells, the basolateral surface of epithelial cells, embryonic mesenchymal cells, vascular smooth muscle cells, endothelial cells and neural cells
30 but not mature circulating B cells. The interaction between CD138 and the PDZ domains of the 41.8kD protein and TIAM1 proteins is believed to be necessary for the proper distribution of CD138 on the cell surface. Disruption of the interaction by administration of an effective

amount of an antagonist is expected to interfere with the migration and adherence of cells to the extracellular matrix, resulting in reduced inflammatory and humoral immune responses. Inhibition of CD138 may be used to treat without limitation diseases such as post-myocardial infarction inflammatory damage, joint injury, rheumatoid arthritis, vasculitis, drug reaction, scleroderma, SLE, Hashimoto thyroiditis, Goodpasture's syndrome, juvenile insulin-dependent diabetes, psoriasis.

E. CD98

As shown *supra*, CD98 interacts with MPP2. CD98 is expressed at high levels on monocytes and at low levels on T cells, B cells, splenocytes, NK cells, and granulocytes. CD98 plays roles in adhesion, fusion and is a L-type amino acid transporter. CD98 is also involved in virus-mediated cell fusion (e.g. paramyoviruses: parainfluenza virus type 2, Newcastle disease virus, and rubulaviruses) and antagonism of CD98 function is expected to treat viral infections and limit viral spread. CD98 inhibitors can be an antiviral agent for, but not limited to paramyovirus, parainfluenza, Newcastle disease and rubula. Other roles include treatment for acute leukemias.

F. CLASP-1

As shown *supra*, CLASP-1 interacts with DLG1, PSD95, and NeDLG. CLASP-1 is a member of a superfamily of immune-cell associated proteins with similar motifs (see PCT/US99/22996 published as WO 00/20434). CLASP-1 functions in the maintenance of the immune synapse. The CLASP-1 transcript is present in lymphoid organs and neural tissue, and the protein is expressed by T and B lymphocytes and macrophages in the MOMA-1 subregion of the marginal zone of the spleen, an area known to be important in T:B cell interaction. CLASP-1 staining of individual T and B cells exhibits a preactivation structural polarity, being organized as a "ball" or "cap" structure in B cells, and forming a "ring", "ball" or "cap" structure in T cells. The placement of these structures is adjacent to the microtubule-organizing center ("MTOC"). CLASP-1 antibody staining indicates that CLASP-1 is at the interface of T-B cell conjugates that are fully committed to differentiation. Antibodies to the extracellular domain of CLASP-1 also block T-B cell conjugate formation and T cell activation.

Antagonism of CLASP-1 function is expected to interfere with immune responses (e.g., T and B cell activation), signal transduction, cell-cell interactions, and

membrane organization. Diseases to be treatment by CLASP-1 agonists/antagonists include, but is not limited to, rheumaotoid arthritis, juvenile diabetes, organ rejection, graft-versus-host disease, scleroderma, multiple sclerosis.

5 G. CLASP-4

As shown supra, CLASP-4 interacts with DLG1, PSD95, NeDLG, LDP, AF6, 41.8, and MINT1. CLASP-4 is a member of a superfamily of immune-cell associated proteins with similar motifs (see copending U.S.Pat. Application 60/196,527 filed April 11, 2000). The CLASP-4 protein is expressed primarily in peripheral blood lymphocytes.

- 10 Inhibition of the interaction of CLASP-4 and PDZ domains will interfere with immune responses (e.g., T and B cell activation), signal transduction, cell-cell interactions, and membrane organization. Disease to be treated bu CLASP-4 agonists/antagonists include, but is not limited to, rheumaotoid arthritis, juvenile diabetes, organ rejection, graft-versus-host disease, scleroderma, multiple sclerosis, acute leukemias, leukemic blast crisis, post-infarction
15 inflammation (cardiac, etc.), atherosclerosis.

H. VCAM1

- The vascular cell adhesion molecule-1 (VCAM-1, CD106) is predominantly expressed by vascular endothelium (i.e., endothelial cells) but has been detected in
20 macrophages, dendritic cells, bone marrow-derived cells, fibroblasts, cortical thymic epithelial cells, vascular smooth muscle cells, myoblasts and myotubes. VCAM-1 mediates adhesion through interacting with an integrin ligand, VLA-4, which is expressed by lymphocytes, monocytes and eosinophils. The interaction between VCAM-1 and VLA-4 is important for activation, flattening and extravasation of VLA-4 expressing cells when the endothelium itself
25 has become activated due to inflammation or injury (Salomon et al., 1997, *Blood* 89:2461-2471; St-Pierre et al., 1996, *Eur. J. Immunol.* 26:2050-2055; Bell et al., 1995, *Int. Immunol.* 7:1861-1871).

- As discussed *supra*, the C-terminal region of VCAM-1 is a ligand for the PDZ domains of MPP1, DLG1, NeDLG1, LDP, 41.8 protein, TIAM1, K807, WWP3 and K303.
30 These interactions are believed to mediate the function of endothelial cell interactions with integrin expressing leukocytes. When the PDZ-PL interactions are disrupted, the adherence of leukocytes to the endothelium will similarly be disrupted, resulting in, e.g., reduction of

inflammation. Thus, inhibition of VCAM-1 binding to PDZ proteins is useful for reducing abnormal VCAM-1 inflammatory responses and associated pathologies such as (but not limited to) renal allograft rejection, insulin-dependent diabetes, rheumatoid arthritis, post-myocardial infarction complications and systemic lupus erythematosus (Pasloske et al., 1994, *Ann Rev Med* 45:283; Ockenhouse et al., 1992, *J. Exp. Med.* 176:1183; Solezk et al., 1997, *Kidney Int.* 51:1476; Tedla, et al., 1999, *Clin. Exp. Immunol.* 117:92-99; Kusterer et al., 1999, *Exp Clin Endocrinol Diabetes* 107:S102-107; Bonomini et al., 1998, *Nephron* 79:399; Suassuna et al., 1994, *Kidney Int.* 46:443; Ferri et al., 1999, *Hypertension* 34:568).

I. CLASP-2

As shown *supra*, CLASP-2 interacts with PSD-95, NeDLG, and DLG1. CLASP-2 is a member of a superfamily of immune-cell associated proteins with similar motifs (see copending U.S. Pat. 09/547,276 filed April 11, 2000; WO 00/10158 filed April 11, 2000; WO 00/10156 filed April 11, 2000). The CLASP-2 transcript is present most strongly in placenta followed by lung, kidney and heart and the protein is expressed in T and B cells, and kidney epithelial cells.

Inhibition of the interaction of CLASP-2 and PDZ domains will interfere with CLASP-2 function resulting in interference with T and B cell function (e.g., T and B cell activation), signal transduction, cell-cell interactions, and membrane organization. In addition, since CLASP-2 is present in heart, blocking CLASP-2 function or expression can selectively block immune responses in the heart (for example, to selectively stop immune response in the heart compartment, e.g., following cardiac transplant rejection or post-MI inflammation, without compromising immunity elsewhere). Other diseases to be treatment by CLASP-1 agonists/antagonists include, but is not limited to, rheumaotoid arthritis, juvenile diabetes, organ rejection, graft-versus-host disease, scleroderma, multiple sclerosis.

J. CD95 (Apo-1/Fas)

CD95 (Fas/Apo-1) and Fas ligand (FasL) are a receptor-ligand pair involved in lymphocyte homeostasis and peripheral tolerance. Binding of Fas by its ligand results in apoptotic cell death, an important major mechanism for safe clearance of unwanted cell during resolution of the acute inflammatory response. As is shown *supra*, CD95 binds the PDZ domains DLG1, PSD95, NeDLG, TIP1, and 41.8. Agents that modulate (e.g., inhibit) the

interaction of CD95 and PDZ domains are useful for treatment of diseases, e.g., organ transplantation, graft-versus-host disease, Crohn's Disease, Ulcerative colitis, inflammatory bowel disease, rheumatoid arthritis, osteoarthritis, multiple sclerosis, scleroderma, mixed connective tissue disease, leukemia and other malignancies.

5

K. KV1.3 (Shaker Type Kv1.3 Potassium Channel)

As shown *supra*, Kv1.3 binds DLG1, PSD95, NeDLG, LIMK, 41.8, RGS12, DVL1, and MINT1. Kv1.3 is a Shaker-related channel protein that is involved in modulating the membrane potential of T lymphocytes (Lewis and Cahalan, 1995, *Ann. Rev. Immunol.* 13:623). Inhibition of the Kv1.3 channel chronically depolarizes the T cell membrane, reduces calcium entry via calcium-activated release calcium channels in the plasma membrane, and consequently inhibits the calcium-signaling pathway essential for lymphocyte activation. Hanada et al., reported that Kv1.3 is associated with DLG1 and PSD95 in Jurkat T cells (*J. Biol. Chem.* 1997, 272:26899). Administration of Kv1.3-PDZ protein agonist/antagonists will

10 13:623). Inhibition of the Kv1.3 channel chronically depolarizes the T cell membrane, reduces calcium entry via calcium-activated release calcium channels in the plasma membrane, and consequently inhibits the calcium-signaling pathway essential for lymphocyte activation. Hanada et al., reported that Kv1.3 is associated with DLG1 and PSD95 in Jurkat T cells (*J. Biol. Chem.* 1997, 272:26899). Administration of Kv1.3-PDZ protein agonist/antagonists will

15 disrupt T cell signaling and can be a useful therapeutic drug to treat, but not limited to, organ transplantation, graft-versus-host disease, Crohn's Disease, Ulcerative colitis, inflammatory bowel disease, rheumatoid arthritis, osteoarthritis, multiple sclerosis, scleroderma, mixed connective tissue disease.

20 L. DNAM-1

As shown *supra*, DNAM-1 binds several PDZ proteins, including MPP1, MPP2, DLG1, PSD95, NeDLG, LIM, AF6, 41.8, RGS12 and WWP3. DNAM-1 is associated with Fyn constitutively but required the presence of pervanadate (a tyrosine phosphatase inhibitor) (Shibuya, et al., 1999, *Immunity*. 1:615-623). Upon stimulation with anti-CD3 or cross-linking

25 DNAM-1 with anti-DNAM-1, DNAM-1 is phosphorylated at Ser329 (Shibuya, et al., 1999, *Immunity* 1:1671-75) and associates with LFA-1. Furthermore, Fyn becomes associated with DNAM-1 independent of pervanadate. Fyn phosphorylates DNAM-1 at Tyr322, but does not require Tyr322 to continue binding to DNAM-1.

Since DNAM-1 itself does not have a SH3 binding domain but has a Src phosphorylation site at Tyr322, an adaptor molecule must be present to bridge DNAM-1 and Fyn. DLG1 has been described in the literature to be present in T cells (Hanada, et al. 1997, *J. Biol. Chem.* 272:26899), but does not bind to Fyn. Although PSD95 does not have a SH3

30 phosphorylation site at Tyr322, an adaptor molecule must be present to bridge DNAM-1 and Fyn. DLG1 has been described in the literature to be present in T cells (Hanada, et al. 1997, *J. Biol. Chem.* 272:26899), but does not bind to Fyn. Although PSD95 does not have a SH3

site, several of the other PDZ proteins do have SH3 binding domains including but not limited to NeDLG, RGS12, WWP3 and MPP2, and can fulfill this function. The adaptor PDZ list *supra* describes binding to DNAM-1 through the PDZ domain. Tyr139 is a candidate phosphorylation site to control association of Fyn to DNAM-1 and the adaptor PDZ. In addition, through WWP3, DNAM-1 may complex with beta-catenin, actin and cadherin (Dobrosotskaya and James, 2000, *Biochem. Biophys. Res. Commun.* 270:903-909).

Based on this analysis, inhibition of PDZ association with DNAM-1 using the reagents of the invention will inhibit Fyn association with DNAM-1 and the subsequent Tyr322 phosphorylation and activation of cytotoxic T cells. Disease that can be treated include but are not limited to Crohn's Disease, multiple sclerosis, ulcerative colitis, inflammatory bowel disease, graft-versus-host, juvenile diabetes, and Hashimoto's Disease.

M. CD83 (HB15)

CD83 is a transmembrane glycoprotein, expressed predominantly on activated dendritic cells (DCs), Langerhans cells in the skin, with some weak expression detected on activated peripheral lymphocytes, and interdigitating reticulum cells within the T cell zones of lymphoid organs (Zhou and Tedder, 1995, *J. Immunol.* 154:3821-3835; Zhou et al., 1992, *J. Immunol.* 149:735-742). CD83 is up-regulated de novo upon activation of an immature DCs, and is the major discriminating marker and characteristic for activated, mature DCs (Czerniecki et al., 1997, *J. Immunol.* 159:3823-3837). DCs function as antigen presenting cells (APCs). Upregulation and expression of CD83 thus appears to be required for DCs to mature and function as APCs.

As shown by experiments described *supra*, the CD83 binds to the PDZ domains of DLG1, PSD95, and NeDLG. These interactions between CD83 and PDZ domains, and between CD83 and DLG1, PSD95, and NeDLG are believed to be important for proper distribution and recycling of CD83. Disruption of CD83 and PDZ proteins with agonists and antagonist can be used to treat, but not limited to, psoriasis, cancers, allergies, autoimmune diseases such as multiple sclerosis, system lupus erythematosus.

N. CD44 (phagocytic glycoprotein 1, lymphocyte homing receptor, p85 and HCAM)

CD44 is single pass transmembrane protein that has several different isoforms due to alternative splicing. It has a broad pattern of expression being detected on both

hematopoietic and non-hematopoietic cell types including epithelial, endothelial, mesenchymal and neuronal cells. CD44H is a major isoform that is expressed in lymphoid, myeloid and erythroid cells (reviewed in Barclay et al., 1997, *The Leukocyte Antigen Facts Book*, 2ed, Academic Press). CD44 is a receptor for hyaluronate (HA), which is a constituent of the extracellular matrix (ECM). In the immune system, CD44 functions as an adhesion molecule on the surface of leukocytes and erythrocytes that binds HA polymers in the ECM, and it can also act as a signaling receptor when HA becomes soluble during inflammatory reactions or tissue damage. The cytoplasmic region of CD44 has been shown to bind or be associated with the actin cytoskeleton through interactions with spectrin and members of the ERM (ezrin, radixin, and meosin) family (reviewed in Lesley et al., 1993; Bajorath, 2000, *Proteins* 39:103-111). Additionally, CD44 is associated with the non-receptor tyrosine kinase p56Lck (Taher et al., 1996, *J. Biol. Chem.* 271:2863-2867. CD44 has been shown to be a co-stimulatory molecule with CD3/TCR engagement to activate T cells (reviewed in Aruffo, 1996, *J. Clin. Invest.* 98:2191-2192).

As described *supra* by experiments reported herein, the C-terminus of CD44 is a ligand for the PDZ domain contained in MPP1, prIL-16 and MINT1. It is believed that the interactions of CD44 with PDZ domains, and between CD44 with MPP1, prIL-16 and MINT1 function in maintenance of leukocyte structure and in leukocyte signaling. Thus, when a CD44-PDZ interaction is disrupted, CD44 will fail to transduce proper intracellular signals, and maintain proper distribution of CD44 on the surface, which will prevent adhesion of leukocytes to the endothelium during inflammation and tissue damage. Administration of agonists/antagonists of this interaction will thus result in, but not limited to, reduced inflammatory responses during tissue ischemia and cell lysis (e.g., rhabdomyosis), vascular insufficiencies (e.g. frostbite), psoriasis, eczema, graft-versus-host disease, granuloma annulare, scleroderma.

O. CD97 (CD55)

As discussed *supra*, CD97 binds the PDZ domains of DLG1 and 41.8. CD97 is a 79.7 kD seven-span transmembrane protein expressed on granulocytes and monocytes and at low levels on resting T cells and B cells. Upon T or B cell activation expression levels of CD97 in T cells and B cells increases rapidly (Eichler et al., 1994, *Scand. J. Immunol.* 39, 111-115; Pickl et al., 1995, *Leukocyte Typing V*:1151-1153). When expressed on COS cells, CD97

confers adhesion to lymphocytes and to erythrocytes.

According to the present invention, the interaction of CD97 with DLG1 and the 41.8 kd protein can be altered to interfere with proper membrane distribution of CD97 and/or recycling of CD97. Such modulation will affect CD97 dependent adherence of cells with therapeutic benefit. Without being limited, agonists and antagonists of CD97-PDZ protein interaction can be used to treat rheumatoid arthritis, osteoarthritis, Crohn's Disease, Ulcerative colitis, psoriasis.

P. Glycophorin C (GC)

As is shown *supra*, the c-terminus of Glycophorin C (GC) interacts with the PDZ domains of human DLG, PSD95, NeDLG, MMP2, AF6, 41.8, and MINT1 (with Mint-1 described previously). Glycophorin C is an integral membrane protein expressed in erythroid cells, thymus, stomach, breast, adult and fetal liver, monocytes, T and B cells (Le Van Kim et al., 1989, *J. Biol. Chem.* 264:20407-20414) and is known for its role in human erythrocytes where it interacts with MPP1 and protein 4.1 to regulate the shape, integrity and mechanical stability of red cells (Marfatia et al., 1997, *J. Biol. Chem.* 272:24191-24197; Reid et al., 1987, *Blood* 69:1068-1072.).

Interactions between Glycophorin C and PDZ proteins DLG, PSD95, NeDLG, MMP2, AF6, 41.8, and MINT1 are believed necessary for maintenance of the physical integrity of cells in which they are expressed. Modulation of GC-PDZ interactions will alter with the function of these and can be utilized to treat, but not limited to, polycythemia vera, spherocytosis.

Q. CDw128A (IL8RA)

As is described *supra*, CDW128A binds to the PDZ domains of DLG1 and NeDLG. There are two forms for the IL-8 receptor, IL-8RA (CDw128A) and IL-8RB (CDw128B) both of which are members of the G protein-coupled receptor superfamily and chemokine receptor branch of rhodopsin family. CDw128A and CDw128B both bind IL-8 with the same affinity but only CDw128B, binds three other IL-8-related CXC chemokines: melanoma growth-stimulating activity (GRO/MGSA), neutrophil-activating peptide 2 (NAP-2) and ENA-78. See, e.g., Ahuja and Murphy, 1996, *J Biol Chem* 271:20545-50.

CDw128A is expressed on all granulocytes, a subset of T cells, monocytes,

endothelial cells, keratinocytes, erythrocytes, and melanoma cells. IL-8 induces chemotaxis of neutrophils, basophils, and T lymphocytes and increases neutrophil and monocyte adhesion to endothelial cells. The binding of IL-8 to IL8RA induces a transient increase in intracellular calcium levels, activation of phospholipase D, a respiratory burst of neutrophils and chemotaxis. This pro-inflammatory response is effective in normal immune responses. Inhibitors of CDw128A are useful for treatment of psoriasis, rheumatoid arthritis, polyarthritis, and for control of angiogenesis-dependent disorders such as melanomas and breast cancer.

R. CD3- η (η)

CD3- η is a splice variant of CD3 zeta and a component of the CD3/TCR complex, which is required for antigen recognition, signal transduction and activation of T cells (Weiss and Littman, 1994, *Cell* 76:263-274.). See, Barclay et al., 1997, The Leucocyte antigen facts book, 2nd Ed, Academic Press. As shown by experiments reported herein, the C-terminal region of CD3- η is a ligand for the PDZ domains of MINT1, 41.8 protein, DLG1, and PSD95.

The interactions of CD3- η with PDZ domains MINT1, 41.8 protein, DLG1, K807, TIP, and PSD95 are believed to be important activation of T cells, which is required for all cellular immune responses. Modulation of this interaction by agonists and antagonists can be used to treat, but is not limited to, acute and chronic allograft rejection, multiple sclerosis, graft-versus-host disease, rheumatoid arthritis.

S. LPAP (CD45-AP, LSM-1)

LPAP is a transmembrane protein expressed on resting T-and B-cells. LPAP has been shown to bind to CD45, a protein that is part of the T-cell receptor complex and has been found to co-localize with CD4, CD2 and Thy-1. LPAP has also been co-immune precipitated with p56(lck) and ZAP-70. The actual function of LPAP is unknown, but data obtained from LPAP deficient mice and Jurkat cell lines suggest that LPAP is an assembly molecule important for the organization of a functional CD45 complex.

As shown *supra*, LPAP binds to DLG-1, MINT-1, KIAA0807 and TIP-1 (sometimes "TAX Interacting Protein 1" or "TAX Interacting Protein" or "TAX IP-1").

Notably, DLG-1, MINT-1, KIAA0807 and TIP-1 are expressed in T-cells. It has been shown that DLG-1 coprecipitates with p56(lck) in T-cells (Hanada et al., 1997, *J. Biol. Chem.* 272(43):26899-26904). The assay described herein also demonstrates that DLG-1 binds



to CD95 and KV1.3, MINT-1 binds to KV1.3, TIP-1 binds to CD95, KV1.3, CD3 η and HTLV-1 TAX oncoprotein. All these molecules are involved in signaling by the TCR or in the regulation of cell death by apoptosis. LPAP is believed to function in organizing the signaling of CD45 in T-cells by recruiting p56(LCD) as a substrate for CD45. Blocking the function of CD45 has been shown to severely impair the T-cell response. The human PDZ domain containing protein KIAA0807 shows a high degree of similarity to the mouse protein MAST205, a serine/threonine protein kinase, and computer based protein domain homology search predicts protein tyrosine kinase activity for KIAA0807. Binding of KIAA0807 to LPAP appears to be involved in, and possibly crucial for, recruiting KIAA0807 encoded protein into the CD45 complex. Further, protein kinase activity of the KIAA0807 protein may add specific kinase function to the CD45 complex and might tune phosphatase activity of CD45. Thus, inhibition of the PDZ domain-PDZ ligand mediated interaction between LPAP and KIAA0807 can be used to alter (e.g., diminish or abolish) immune response.

The assay described herein also shows that TIP-1 binds to LPAP and to CD95. CDp5 has been shown to be a pivotal component of programmed cell death (apoptosis). Binding of TIP-1 to CD95 might be regulated by changes in phosphorylation of either binding partner, and LPAP mediated recruitment of TIP-1 into the CD45 (phosphatase / kinase)-complex might ensure proximity of phosphatase / kinase activity when TIP-1 engages in CD95 binding. In addition, TIP-1 / LPAP binding might compete for TIP-1 / CD95 binding, thus being involved in switching from T-cell proliferation to apoptosis.

Studies with LPAP null mice have shown impaired lymphocyte responses to antigen receptor (Motoya, et al., 1999, *J. Biol. Chem.* 274(3):1407-1414) and have suggested that LPAP plays a role in regulation of lymphocyte expansion in particular lymphatic organs (Ding, et al., 1999, *Eur. J. Immunol.* 29(12):3956-3961). Therefore, inhibiting the interaction between LPAP and PDZ proteins is expected to alter the CD45-mediated path from the rest of the immune response and to change the pattern of lymphocyte distribution. Agonists and antagonist of PL – PDZ binding can be used to treat a variety of disease, including immune disorders such as (but not limited to) rheumatoid arthritis, transplant rejection, multiple sclerosis, scleroderma, graft-versus host disease.

T. CD46 (Complement membrane cofactor protein (MCP))

CD46 is a membrane protein expressed on all nucleated cells, but not on

erythrocytes. CD46 is a member of the regulator of complement activation protein family. Its primary function is the protection of cells from complement attack by inactivating membrane deposited C3b/C4b complement (Liszewski, et al., 1999, *Adv. Immunol.* 61:201-283). CD46 exists in more than 8 isoforms that are generated by differential splicing, with molecular weights ranging from 45 to 70 kD. In addition to the above function, CD46 also serves as the receptor for the measles virus and for other pathogenic microorganisms (e.g. *Streptococcus pyogenes*) (Manchester et al, 1994, *Proc. Natl Acad. Sci. USA* 91:2161; Okada et al., 1995, *Proc. Natl Acad. Sci. USA* 92:2489-2493.). CD46 also appears to be over-expressed on certain tumors (Jurianz et al., 1999, *Mol Immunol* 36:929-939) thus rendering tumor cells insensitive to the action of complement. See, Barclay et al., (1997) *The Leucocyte antigen facts book*, 2nd ed, Academic Press.

As shown *supra*, CD46 binds DLG1, PSD95 and Ne-DLG. This interaction is believed to be necessary for proper membrane distribution of CD46 and/or recycling of CD46. Alteration of the CD46-PDZ interaction can reduce the ability of measles virus and other pathogens to enter cells, renders CD46-expressing tumors susceptible to attack by complement. The administration of CD46-PDZ interaction agonists and antagonists is useful for the treatment of , but not limited to, cancers and viral infectious diseases.

U. CDw128B

As is described *supra*, CDw128B binds to the PDZ domains of DLG1, NeDLG, PSD95, K807 and 41.8 in the assays described *supra*. There are two forms for the IL-8 receptor, IL-8RA (CDw128A) and IL-8RB (CDw128B) both of which are members of the G protein-coupled receptor superfamily and chemokine receptor branch of rhodopsin family. CDw128A and CDw128B both binds IL-8 with equal affinity but only CDw128B, also binds three other IL-8-related CXC chemokines: melanoma growth-stimulating activity (GRO/MGSA), neutrophil-activating peptide 2 (NAP-2) and ENA-78. See, e.g., Ahuja, SK and Murphy, PM. 1996. *J Biol Chem* 271:20545-50.

CDw128B is expressed on all granulocytes, a subset of T cells, monocytes, endothelial cells, keratinocytes, erythrocytes, and melanoma cells. IL-8 induces chemotaxis of neutrophils, basophils, and T lymphocytes but diminished relative to IL8RA and increases neutrophil and monocyte adhesion to endothelial cells. The binding of IL-8A to its receptor induces a transient increase in intracellular calcium levels and granule release but does not

induce activation of phospholipase D or a respiratory burst in neutrophils. This pro-inflammatory response is effective in normal immune responses. Inhibitors of CDw128B are useful for treatment of psoriasis, rheumatoid arthritis, polyarthritis, and for control of angiogenesis-dependent disorders such as melanomas and breast cancer.

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V. DOCK2

The DOCK family is a group of transmembrane proteins that interact with the cytoskeleton to affect cell shape, and maintain structural integrity of functional subdomains within a cell. Members of this new family include *Drosophila myoblast city* (mbc), DOCK180 (DOCK1), DOCK2, DOCK3, CED5, KIAA0209 and CLASP. The prototypical molecule, DOCK1 or DOCK180 is the human homologue of the *C. elegans* gene, CED5, which is involved in the engulfment and phagocytosis by macrophages of cells undergoing apoptosis (apoptotic cells). Of the human family members, DOCK2 is most closely related to DOCK1. DOCK2 expression appears to be confined to cells of leukocytic origin. DOCK2 is found in peripheral blood lymphocytes and can convert a flatten cell into a rounded morphology upon transfection (Nagase, et. al., 1996, *DNA Res* 3:321-29; Nishihara, 1999, *Hokkaido Igaku Zasshi* 74:157). DOCK2 mRNA is highly expressed in peripheral blood lymphocytes, with lower expression in thymus and spleen and very weak expression in colon and small intestine. Immunohistochemistry with antibodies directed against DOCK2 detect expression of DOCK2 in macrophages in the interstitium and alveoli. DOCK2 protein is also detected in lymphocytes of the lymph node, and macrophage and lymphocytes in tonsillar tissue (Nishihara et al. 1999. *Bioch. Biophys. Acta* 1452:179). DOCK2, like DOCK1, has been shown to bind Rac1, a GTPase, which could account for it's ability to round-up flat NRK (normal rat kidney) cells (Nishihara, 1999, *Hokkaido Igaku Zasshi* 74:157).

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As shown supra, DOCK2 is a PL and binds to the PDZ-containing proteins KIAA807, DLG1, PSD95, NeDLG, Syntrophin α 1 (Synt α 1) and KIAA0561. In addition to a PDZ domain, KIAA807 contains a kinase domain suggesting that the interaction between KIAA807 and DOCK may have a role in signal transduction. The function of KIAA0561 is unknown at this time. DLG1, PSD95, NeDLG and Syntrophin α 1 have roles in cell signaling and structure (Gomperts, 1996, *Cell* 84:659; Shen and Wyszynski, 1997, *Bioessays* 19:847; Sheng and Kim, 1996, *Curr. Opin. Neurobiol.* 6:602; Leu et al., 1994, *Proc. Nat. Acad. Sci.* 91:9818). Most notably, PSD95, NeDLG and Syntrophin α 1 are necessary for proper

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organization of neuronal synapses (Adams et al., 2000, *J. Cell Biol.* 150:1385; Masuko et al., 1999, *J. Biol. Chem.* 274:5782). Using PDZ adaptor proteins, DOCK2 would be required to control cell shape of lymphocytes for their transit through the vascular circulation through its direct or indirect interaction with Rac1. The PDZ-PL interactions would be necessary for proper assembly and maintenance of the immunological synapse in a similar manner for their role in the neuronal synapse. These interactions would allow proper cell-to-cell interactions required to initiate immune reactions. Modulation of DOCK2-PDZ interactions by agonists and antagonists can be used to treat diseases such as, but not limited to, acute lymphocytic leukemias, leukemic blast crisis, post-myocardial infarction inflammation, and post-traumatic inflammation.

W. CD34

As is shown *supra*, CD34 binds DLG1, PSD95, K807, and NeDLG. CD34 is expressed on a small subpopulation of bone marrow cells which includes hematopoietic stem cells. CD34 is also present on bone marrow stromal cells and on endothelial cells. The selectins CD62L (L-selectin) and CD62E (E-selectin) bind CD34. CD34 mediates attachment and rolling of leukocytes. The hematopoietic stem cell properties of CD34 include myeloid differentiation of stem cells. Modulation of the CD34-PDZ interaction with agonists and/or antagonists can be used to treat, but is not limited to, myelodysplasia, leukemias, post-traumatic inflammation, post-myocardial infarction inflammation.

X. Fc epsilon receptor beta I chain (FcεRβI)

The high affinity receptor for human IgE, FcεRI, is composed of an α, β, and disulfide-linked γ homodimer. The α-chain binds the Fc portion of IgE, whereas the β-chain serves to amplify signals that are transduced through the γ-chain homodimer. Both αβγ2 tetramer and αγ2 trimer complexes exist, but the β-chain amplifies the signal 5- to 7-fold, as measured by Syk activation and calcium mobilization. Additionally, the FcεRβI is a PDZ ligand and is a member of the CD20/ FcεRβI receptor family. As is shown *supra*, FcεRβI binds MINT1.

As the high-affinity receptor for IgE, FcβRI on basophils and mast cells plays a central role in the initiation of allergic responses. Signaling through the FcβRI begins by crosslinking of a multivalent allergen bound to IgE. The result is vesicular degranulation,

release of histamine, leukotrienes and pro-inflammatory cytokines (IL-6 and TNF α), factors responsible for the symptoms of immediate hypersensitivity. Alteration of signaling by targeting PL/PDZ interaction with agonists and antagonists can be used to treat, but is not limited to, asthma, atopic dermatitis, eczema, drug reaction, mastocytosis, urticaria, eosinophilia myalgia syndrome (Turner, H., et. al., 1999, *Nature* 402 SUPP:B24).

Y. FAS Ligand (FasL)

CD95 (Fas/Apo-1) and Fas ligand (FasL) are a receptor-ligand pair critically involved in lymphocyte homeostasis and peripheral tolerance. Binding of Fas by its ligand results in apoptotic cell death, an important major mechanism for safe clearance of unwanted cell during resolution of the acute inflammatory response. FasL is mainly restricted to activated T lymphocytes and is rapidly induced. Fas ligand is frequently up-regulated in breast cancer, as compared with normal breast epithelial cells and benign breast disease. As is shown *supra*, FasL binds the KIAA0561 PDZ domain. The PDZ-PL modifiers are useful for treatment of, but not limited to, tumors, e.g., tumors unresponsive to conventional chemotherapy.

Z. CDW125 (IL5R)

As is shown *supra*, CDW125 binds PTN-4 and RGS12. CDW125 is an IL-5 receptor expressed on eosinophils and basophils. IL5 promotes growth and differentiation of eosinophil precursors and activates mature eosinophils (Takatsu et al 1994, *Adv. Immunol.* 57:145-190). The secreted form of CDW125 has antagonistic properties and is able to inhibit IL-5-induced eosinophil proliferation and differentiation. Modulation of CDW125 binding to PDZ domains may be used to treat, but is not limited to, asthma, atopic dermatitis, eczema, drug reaction, urticaria, mastocytosis, eosinophilia.

AA. Burkitt's lymphoma receptor-1 (BLR-1; CXCR5)

BLR-1 is a transmembrane receptor detected primarily on B cells, and shown to be upregulated in stimulated T cells (Dobner et al., 1992, *Eur J. Immunol.* 22:2795-2799.; Flynn et al., 1998, *J. Exp. Med.* 188:297-304). BLR-1 functions in chemotaxis of B and T cells into follicles of secondary lymphoid organs (e.g. spleen) for proper development and selection toward antigens (Forster et al., 1996, *Cell* 87:1037-1047. Its ligand is B-lymphocyte chemoattractant (BLC), which is strongly expressed in the follicles of Peyer's patches, spleen

and lymph nodes (Gunn et al., 1998, *Nature* 391:799-803). Consistent with its chemotactic role is the demonstration that BLR-1 expression is downregulated in developed, activated B cells (plasma cells) to prevent them from being retained in follicles (Forster et al., 1994, *Cell Mol Biol* 40:381-387), and blr ^{-/-} B cells fail to migrate into B cell follicles (Forster et al., 1996).

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It has been unclear how BLR-1 is organized on the cell surface and how the signaling occurs intracellularly. Through the PRISM MATRIX assay, we have deduced several intracellular proteins that interacts with BLR-1 that may play an important role in organizing intracellular and extracellular events related to BLR-1 function in B cell maturation. Disruption of PL/PDZ-mediated interactions between BLR-1 and PDZ domain molecules would be anticipated to interrupt BLR-1 signaling during B cell maturation, with implications for treatment and diagnosis of immune deficiency, allergy and autoimmunity.

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PDZ domain containing proteins function by clustering proteins to organize macromolecular protein complexes, including modulation of protein phosphorylation. As shown *supra*, BLR-1 binds to the PDZ domain containing molecules MINT1, PDZK1, KIAA0807 and KIAA0561. Notably, MINT1, PDZK1, KIAA0807 and KIAA0561 are expressed in B-cells. Without intending to be bound by a particular mechanism, we suggest, that BLR-1 interactions with MINT1 and PDZK1 are crucial for proper BLR-1 clustering into functional macromolecular complexes whereas BLR-1 binding to KIAA0807 and KIAA0561 is likely to be associated with changes in the phosphorylation state of BLR-1 (or other molecules associated with BLR-1). In turn, BLR-1 phosphorylation may regulate BLR-1 binding to PDZK1 and MINT1.

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MINT1 is known as part of a multi-protein complex that occurs in the neurooogical synapse. This complex functions as an intermediate for synaptic vesicle fusion and / or vesicle docking (Okamoto and Sudhof, 1997, *J. Biol. Chem.* 272(50):31459-31464). We suggest, that binding of MINT1 to BLR-1 as detected by the herein described assay might constitute a crucial step in recruitment, transport and/or functional organization of active BLR-1 complexes.

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In several instances, the PDZK1 protein has been demonstrated to be a potent organizer of functional membrane bound multimeric protein complexes. PDZK1 is a four-PDZ domain molecule which is peculiar in that its carboxy terminal end constitutes a PL that has the property to interact with the PDZK1 most N-terminal PDZ domain (Kocher, et al., 1999, *Lab*

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Invest 79:1161-70). MAP17, cMOAT, CFTR, and scavenger receptor type B are four known examples of membrane bound and clustered proteins that engage via PDZ / PL mediated interactions with PDZ domains of PDZK1. MAP17 is expressed in normal kidney epithelium, but it is strongly upregulated in human kidney, colon, lung and breast carcinomas (Kocher, et al., *Lab Invest.*, 1998, 78:117-125). cMOAT (canalicular multispecific anionic transporter) is a protein involved in multidrug resistance and is overexpressed in several carcinoma cells lines; (Kocher, et al., *Lab Invest.*, 1999, 79:1161-1170). CFTR functions as a chloride channel which causes cystic fibrosis when mutated. CFTR binds to PDZK1 PDZ domains 2-4. The clustering of several CFTR molecules mediated by PDZK-1 has been shown to potentiate CFTR function (Wang et al., 2000, *Cell*, 103:169-179). Scavenger receptor type B is expressed in liver cells and is involved in the uptake of cholesterol from high density lipoproteins. The interaction between scavenger receptor type B and PDZK-1 seem to be associated with the regulation of atherogenesis (Ikemoto et al., 2000, *Proc. Natl. Acad. Sci.* 97:6538-6543).

The above described examples demonstrate that proper spatial organization and concentration of MAP17, cMOAT, CFTR, and scavenger receptor type B are vitally correlated with their proper physiological function. Furthermore, these observations strongly suggest that PDZK1 has a pivotal role in recruiting and in organizing these multimeric membrane associated protein complexes. We suggest, that PDZK1 is also crucial for recruiting and organizing BLR-1 into a functional membrane associated complex.

Intra-molecular binding of the PDZK1 PL to N-terminal PDZK1 PDZ domain might ensure proper orientation of up to three PL molecules (homo- or heterotrimer) bound to PDZK1 domains 2, 3, and 4. In addition, PDZK1 PDZ domain 1 might interact with the PL of another PDZK1 molecule in a head-to-tail fashion (inter-molecular), thus enabling the local formation of stable high density homo- or heteromeric complexes. The equilibrium between intra-molecular and inter-molecular PDZK1 PL/PDZ binding might depend on the local PDZK1 concentration, whereas phosphorylation changes at the PDZK1 PL and / or the BLR-PL might function as an on / off switch for either interaction. BLR-1 PL phosphorylation might be regulated by KIAA0807 and KIAA0561 proteins when interacting with BLR-1.

KIAA0807 and KIAA0561 encode human PDZ domain containing proteins that are highly similar to the mouse protein MAST205, a serine/threonine protein kinase, and computer based protein domain homology search predicts protein tyrosine kinase activity for both, KIAA0807 and KIAA0561. PL / PDZ mediated binding of BLR-1 to KIAA0807 and to

KIAA0561 proteins as detected by the herein described assay might change the phosphorylation status of BLR-1 and thus regulate its activity and binding properties; in addition, the BLR-1 / KIAA0807 and the BLR-1/KIAA0561 interactions might associate protein kinase activity with BLR-1 which, in turn, might modulate the phosphorylation status of PDZK1 or any other of its PDZ domain containing interacting proteins in the course of (BLR-1) transport to and / or clustering in the cell membrane. The phosphorylation status of either binding partner might constitute the switch between engagement into and interruption of a PL/PDZ mediated protein-protein interaction.

As shown by experiments reported herein, the C-terminal end of BLR-1 binds to the PDZ domain containing proteins PDZK1, MINT1, KIAA0807 or KIAA0561. Without intending to be bound by a particular mechanism, the interaction between BLR-1 and the PDZ domain proteins is necessary for the proper distribution and signaling of BLR-1 on the cell surface. Normal function of the receptor is required for physiological B lymphocyte chemotaxis and thus for B lymphocyte function. This function of the receptor is disrupted by small molecule therapeutics that disrupt BLR-1/PDZ binding (see, e.g., Example 7, *infra*). When this interaction is disrupted, the chemotactic abilities of lymphoid cells expressing BLR-1 is similarly disrupted. Such a disruption results in a reduced immune response, interference with the ability of lymphocytes to properly circulate and develop responses to antigen. Such small molecule therapeutics are thus immunosuppressive agents that specifically target B lymphocytes. These therapeutics are of use in the treatment of autoimmune disorders involving overactivity of B lymphocytes, such as rheumatoid arthritis, systemic lupus erythematosus, and pemphigus vulgaris. Agonists and antagonists of the BLR-PDZ interaction are used to treat immune system diseases including, but is not limited to, rheumatoid arthritis, transplant rejection, multiple sclerosis, scleroderma, graft-versus host disease, systemic lupus erythematosus, scleroderma and other autoimmune diseases.

BB. CD4

CD4 is a co-receptor with the T cell receptor (TCR) involved in antigen recognition. Both CD4 and TCR belong to the immunoglobulin supergene family. T cell activation is enhanced by increasing the avidity of T cells for effector and target cells. The cytoplasmic domain is involved in signal transduction and association with the tyrosine kinase p56^{lck}. CD4 is expressed on most thymocytes, two-thirds of peripheral blood T lymphocytes,

The logo of the American Society of Human Genetics (ASHG) is a circular emblem. It features a central design with the text "AMERICAN SOCIETY OF HUMAN GENETICS" around the perimeter. The center contains a stylized representation of a human figure or a genetic structure, possibly a chromosome or a DNA helix, with the words "HUMAN GENETICS" integrated into the design.

[illegible]

CC. PAG (Phosphoprotein Associated With GlycoSphingolipid-Enriched Microdomains)

PAG is a recently identified, transmembrane adaptor phosphoprotein. It is expressed in hematopoietic cells including peripheral blood lymphocytes, monocytes and neutrophils, and is a substrate for kinases including the Lck and Fyn (Brdicka et al. 2000 *J. Exp. Med.* 191:1592-1604). We have discovered using the methods of the invention (see Table 2) that PAG contains a PDZ-ligand motif and have demonstrated that PAG binds the PDZ-containing protein kinase KIAA807. This PDZ-PL interaction is consistent with an adaptor or scaffolding role for PAG.

Subcellularly, PAG is found with glycosphingolipid-enriched microdomains (GEMs, also known as lipid rafts or detergent-insoluble glycolipid-rich membrane domains). GEMs are 70nm detergent resistant membrane islands found within the bulk plasma membrane of a cell. Each GEM has a concentration of lipids with higher saturated fatty acid side-chains, which favors their association. Importantly, GEMs also contain vital signal transduction molecules including kinases (e.g. Lck, Fyn, LAT, and PI-3'-kinase), adaptor proteins (LAT) and G-proteins. In hematopoietic cells GEMs are a required, functional component of immune cell activation since disruption of GEMs attenuates their activation. Additionally, GEMs functionally interact with the lymphocyte cytoskeleton, which is necessary for microtubule organization and proper lymphocyte activation (Xavier and Seed, 1999, *Curr. Op. in Immunol.* 11:265-269; Xavier et al., 1998, *Immunity* 8:723-732; Montixi et al., 1998, *EMBO J.* 17:5334-5348).

The identification of PAG as a component of GEMs and a target of kinases found within GEMs indicates that PAG is an important mediator of cellular signal transduction, and is able to recruit other signaling components, namely KIAA807, to GEMs and regulate signal transduction. PAG has been shown to be a negative regulator of T-cell activation (Brdicka et al., 2000, *J. Exp. Med.* 191:1592-1604), and the kinase activity of KIAA807 may transduce the negative regulatory effects of PAG. Such an association provides a means to modulate immune function. As a negative regulator of T-cell activation, enhancement of the PAG:KIAA807 interaction may reduce an overactive immune system in autoimmune disease states during transplantation rejection, rheumatoid arthritis, systemic lupus erythematosus (SLE) and multiple sclerosis. In contrast disruption of this interaction may stimulate the activation of lymphocytes, which could help patients with immune deficiencies such as HIV-induced AIDS.

6.5 Agonists and Antagonists of PDZ-PL Interactions

As described herein, interactions between PDZ proteins and PL proteins in cells (e.g., hematopoietic cells, e.g., T cells and B cells) may be disrupted or inhibited by the administration of inhibitors or antagonists. Inhibitors can be identified using screening assays described herein. In embodiment, the motifs disclosed herein are used to design inhibitors. In some embodiments, the antagonists of the invention have a structure (e.g., peptide sequence) based on the C-terminal residues of PL-domain proteins listed in TABLE 2. In some embodiments, the antagonists of the invention have a structure (e.g., peptide sequence) based on a PL motif disclosed herein.

The PDZ/PL antagonists and antagonists of the invention may be any of a large variety of compounds, both naturally occurring and synthetic, organic and inorganic, and including polymers (e.g., oligopeptides, polypeptides, oligonucleotides, and polynucleotides), small molecules, antibodies, sugars, fatty acids, nucleotides and nucleotide analogs, analogs of naturally occurring structures (e.g., peptide mimetics, nucleic acid analogs, and the like), and numerous other compounds. Although, for convenience, the present discussion primarily refers to antagonists of PDZ-PL interactions, it will be recognized that PDZ-PL interaction agonists can also be used in the methods disclosed herein.

In one aspect, the peptides and peptide mimetics or analogues of the invention contain an amino acid sequence that binds a PDZ domain in hematopoietic cells such as T cells and B cells, or otherwise inhibits the association of PL proteins and PDZ proteins. In one embodiment, the antagonists comprise a peptide that has a sequence corresponding to the carboxy-terminal sequence of a PL protein listed in TABLE 2, e.g., a peptide listed in TABLE 4. Typically, the peptide comprises at least the C-terminal two (2), three (3) or four (4) residues of the PL protein, and often the inhibitory peptide comprises more than four residues (e.g., at least five, six, seven, eight, nine, ten, twelve or fifteen residues) from the PL protein C-terminus. See, e.g. Section 6.5.1, *infra*. Moreover, the C-terminal domains of specific surface receptors expressed by hematopoietic system and endothelial cells may themselves be used as inhibitors, and may be used as the basis for rational design of non-peptide inhibitors. See Section 6.6, *infra*.

In some embodiments, the inhibitor is a peptide, e.g., having a sequence of a PL C-terminal protein sequence. See, e.g. Section 6.5.1, *infra*.

In some embodiments, the antagonist is a fusion protein comprising such a sequence. Fusion proteins containing a transmembrane transporter amino acid sequence are particularly useful. See, e.g. Section 6.9, *infra*.

5 In some embodiments, the inhibitor is conserved variant of the PL C-terminal protein sequence having inhibitory activity. See, e.g. Section 6.5.2, *infra*.

In some embodiments, the antagonist is a peptide mimetic of a PL C-terminal sequence. See, e.g. Section 6.5.3, *infra*.

In some embodiments, the inhibitor is a small molecule (i.e., having a molecular weight less than 1 kD). See, e.g. Section 6.5.4, *infra*.

10 6.5.1 Peptide Antagonists

In one embodiment, the antagonists comprise a peptide that has a sequence of a PL protein carboxy-terminus listed in TABLE 2. The peptide comprises at least the C-terminal two (2) residues of the PL protein, and typically, the inhibitory peptide comprises more than two residues (e.g., at least three, four, five, six, seven, eight, nine, ten, twelve or fifteen residues) from the PL protein C-terminus. The peptide may be any of a variety of lengths (e.g.,
15 at least 2, at least 3, at least 4, at least 5, at least 6, at least 8, at least 10, or at least 20 residues) and may contain additional residues not from the PL protein. It will be recognized that short PL peptides are sometime used in the rational design of other small molecules with similar properties.

20 Although most often, the residues shared by the inhibitory peptide with the PL protein are found at the C-terminus of the peptide. However, in some embodiments, the sequence is internal. Similarly, in some cases, the inhibitory peptide comprises residues from a PL sequence that is near, but not at the c-terminus of a PL protein (see, Gee et al., 1998, *J Biological Chem.* 273:21980-87).

25 Sometime the PL protein carboxy-terminus sequence is referred to as the “core PDZ motif sequence” referring to the ability of the short sequence to interact with the PDZ domain. For example, in an embodiment, the “core PDZ motif sequence” of a hematopoietic cell surface receptor at its C-terminus contains the last four amino acids, this sequence may be used to target PDZ domains in hematopoietic cells. As described above, the four amino acid
30 core of a PDZ motif sequence may contain additional amino acids at its amino terminus to further increase its binding affinity and/or stability. Thus, in one embodiment, the PDZ motif

sequence peptide can be from four amino acids up to 15 amino acids. It is preferred that the length of the sequence to be 6-10 amino acids. More preferably, the PDZ motif sequence contains 8 amino acids. Additional amino acids at the amino terminal end of the core sequence may be derived from the natural sequence in each hematopoietic cell surface receptor or a synthetic linker. The additional amino acids may also be conservatively substituted. When the third residue from the C-terminus is S, T or Y, this residue may be phosphorylated prior to the use of the peptide.

In some embodiments, the peptide and nonpeptide inhibitors of the are small, e.g., fewer than ten amino acid residues in length if a peptide. Further, it is reported that a limited number of ligand amino acids directly contact the PDZ domain (generally less than eight) (Kozlov et al., 2000, Biochemistry 39, 2572; Doyle et al., 1996, Cell 85, 1067) and that peptides as short as the C-terminal three amino acids often retain similar binding properties to longer (> 15) amino acids peptides (Yanagisawa et al., 1997, J. Biol. Chem. 272, 8539).

FIGURES 3A-H show the use of peptides to inhibit PL-PDZ interactions using the G assay described *supra*. In **FIGURE 3A and B**, the inhibition assays were carried out using GST fusion proteins containing PDZ domains from DLG1 or PSD95 (see *supra* and **TABLE 3**). Binding of biotinylated PL peptides for Clasp 2, CD46, Fas, or KV1.3 (as listed in **TABLE 4**) was determined in the presence of various competitor peptides (at a concentration of 100 uM) or in the absence of a competitor (equalized as 100% binding). The competitor peptides were 8-mers peptides having the sequence of C-terminus of Clasp 2 (MTSSSSVV, SEQ ID NO: 191), CD46 (REVKFTSL, SEQ ID NO: 113), or Fas (RNEIQSLV, SEQ ID NO: 48), a unlabeled 19-mer having the sequence of c-terminus of KV1.3 (i.e., non-biotinylated AA33L as listed in **TABLE 3**), or a peptide having the sequence of residues 64-76 of hemoglobin (Vidal et al., 1999, *J. Immunol.* 163, 4811), i.e., an unrelated competitor. The binding of biotinylated peptide (10 uM for Fas and KV1.3, 20 uM for Clasp 2 and CD46) to GST alone was subtracted from the binding to the fusion proteins to obtain the net signal for each experimental condition. This net signal was then normalized by dividing by the signal in the absence of competitor peptide and the data were plotted. Error bars indicated the standard deviation of duplicate measurements. Specific inhibition of Clasp 2 PL-DLG PDZ binding was observed with the CLASP-2 8-mer, the CD46 8-mer, the FAS-8-mer, and the KV13 peptide, but not in the absence of peptide or using an unrelated peptide.

FIGURES 3C-F show similar assays using shorter peptides to inhibit (e.g., a

3-mer and a 5-mer). Figures 3C-E show binding of biotinylated PL peptides for Clasp 2, CD46, Fas, or KV1.3, at the indicated concentration (as listed in **TABLE 3**) to GST fusion proteins containing PDZ domains from NeDLG, DLG1, or PSD95 in the absence or presence of 1 mM 3-mer peptide having the sequence of the C-terminus of Clasp 2 (SVV). (Table 3). **FIGURE 3F** shows the effect on binding of a 5-mer CD49E peptide (ATSDA, SEQ ID NO: 25) to GST fusion proteins containing a PDZ domain from 41.8Kd

6.5.2 Peptide Variants

Having identified PDZ binding peptides and PDZ-PL interaction inhibitory sequences, variations of these sequences can be made and the resulting peptide variants can be tested for PDZ domain binding or PDZ-PL inhibitory activity. In embodiments, the variants have the same or a different ability to bind a PDZ domain as the parent peptide. Typically, such amino acid substitutions are conservative, i.e., the amino acid residues are replaced with other amino acid residues having physical and/or chemical properties similar to the residues they are replacing. Preferably, conservative amino acid substitutions are those wherein an amino acid is replaced with another amino acid encompassed within the same designated class.

6.5.3 Peptide Mimetics

Having identified PDZ binding peptides and PDZ-PL interaction inhibitory sequences, peptide mimetics can be prepared using routine methods, and the inhibitory activity of the mimetics can be confirmed using the assays of the invention. Thus, in some embodiments, the antagonist is a peptide mimetic of a PL C-terminal sequence. The skilled artisan will recognize that individual synthetic residues and polypeptides incorporating mimetics can be synthesized using a variety of procedures and methodologies, which are well described in the scientific and patent literature, e.g., Organic Syntheses Collective Volumes, Gilman et al. (Eds) John Wiley & Sons, Inc., NY. Polypeptides incorporating mimetics can also be made using solid phase synthetic procedures, as described, e.g., by Di Marchi, et al., U.S. Pat. No. 5,422,426. Mimetics of the invention can also be synthesized using combinatorial methodologies. Various techniques for generation of peptide and peptidomimetic libraries are well known, and include, e.g., multipin, tea bag, and split-couple-mix techniques; see, e.g., al-Obeidi (1998) Mol. Biotechnol. 9:205-223; Hruby

(1997) Curr. Opin. Chem. Biol. 1:114-119; Ostergaard (1997) Mol. Divers. 3:17-27; Ostresh (1996) Methods Enzymol. 267:220-234.

6.5.4 Small Molecules

In some embodiments, the inhibitor is a small molecule (i.e., having a molecular weight less than 1 kD). Methods for screening small molecules are well known in the art and include those described *supra* at Section 6.4.

6.6 Cell Surface Receptors and PDZ-Domain Binding Sequences

The following sections describe specific surface receptors expressed by different cell types in the hematopoietic and immune response system. The C-termini of these receptors are used as inhibitors, or serve as the basis for designing PDZ motif sequence peptides, variants, fusion proteins, peptidomimetics, and small molecules for use in inhibiting PDZ-PL interactions. In a preferred embodiment, the peptides are tested in an assay of the invention for inhibitory or modulatory activity (also see, TABLE 4, and discussion *supra*).

6.6.1 PDZ Motif Sequences of T Cell Surface Receptors

A number of surface receptors expressed by T cells contain a PDZ motif sequence (PL sequence). These molecules include CD3 η , CD4, CD6, CD38, CD49e, CD49f, CD53, CD83, CD90, CD95, CD97, CD98, CDw137 (41BB), CD166, CDw128 (IL8 R), DNAM-1, Fas ligand (FasL) and LPAP (Barclay et al., 1997, The Leucocyte Antigen Facts Book, second edition, Academic Press), CLASP-1, CLASP-2, CLASP-4, KV1.3, BLR-1 (CXCR5), PAG and DOCK2.

The C-terminal core sequence of CD3 is SSQL (SEQ ID NO:4). When naturally-occurring residues are added or removed from the core sequence, QL (SEQ ID NO:), SQL (SEQ ID NO:), SSSQL (SEQ ID NO:5), SSSSQL (SEQ ID NO:6), PSSSSQL (SEQ ID NO:7), and PPSSSSQL (SEQ ID NO:8) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD4 is CSPI (SEQ ID NO:9). When naturally-occurring residues are added or removed from the core sequence, PI (SEQ ID NO:), SPI (SEQ ID NO:), TCSPI (SEQ ID NO:10), KTCSPI (SEQ ID NO:11), QKTCSPI (SEQ ID NO:12), and FQKTCSPI (SEQ ID NO:13) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD6 is ISAA (SEQ ID NO:14). When naturally-occurring residues are added or removed from the core sequence, AA (SEQ ID NO:), SAA (SEQ ID NO:), DISAA (SEQ ID NO:15), DDISAA (SEQ ID NO:16), YDDISAA (SEQ ID NO:17), and DYDDISAA (SEQ ID NO:18) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD38 is TSEI (SEQ ID NO:19). When naturally-occurring residues are added or removed from the core sequence, EI (SEQ ID NO:), SEI (SEQ ID NO:), CTSEI (SEQ ID NO:20), SCTSEI (SEQ ID NO:21), SSCTSEI (SEQ ID NO:22), and DSSCTSEI (SEQ ID NO:23) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD49e is TSDA (SEQ ID NO:24). When naturally-occurring residues are added or removed from the core sequence, DA (SEQ ID NO:), SDA (SEQ ID NO:), ATSDA (SEQ ID NO:25), PATSDA (SEQ ID NO:26), PPATSDA (SEQ ID NO:27), and KPPATSDA (SEQ ID NO:28) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD49f is TSDA (SEQ ID NO:29). When naturally-occurring residues are added or removed from the core sequence, DA (SEQ ID NO:), SDA (SEQ ID NO:), LTSDA (SEQ ID NO:30), RLTS DA (SEQ ID NO:31), ERLTSDA (SEQ ID NO:32), and KERLTSDA (SEQ ID NO:33) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD53 is TIGL (SEQ ID NO:34). When naturally-occurring residues are added or removed from the core sequence, GL (SEQ ID NO:), IGL (SEQ ID NO:), QTIGL (SEQ ID NO:35), SQTIGL (SEQ ID NO:36), TSQTIGL (SEQ ID NO:37), and KTSQTIGL (SEQ ID NO:38) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD83 is TELV (SEQ. ID. NO: 177). When naturally-occurring residues are added or removed from the core sequence, LV (SEQ ID NO:), ELV (SEQ ID NO:), KTELV (SEQ. ID. NO: 178), HKTELV (SEQ. ID. NO: 179), PHKTELV (SEQ. ID. NO: 180), and TPHKTELV (SEQ. ID. NO: 181) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD90 is FMSL (SEQ ID NO:39). When naturally-occurring residues are added or removed from the core sequence, SL (SEQ ID NO:), MSL (SEQ ID NO:), DFMSL (SEQ ID NO:40), TDFMSL (SEQ ID NO:41), ATDFMSL (SEQ ID NO:42), and QATDFMSL (SEQ ID NO:43) may also be used to target a PDZ domain-containing protein in T cells.



The C-terminal core sequence of CD95 is QSLV (SEQ ID NO:44). When naturally-occurring residues are added or removed from the core sequence, LV (SEQ ID NO:), SLV (SEQ ID NO:), IQSLV (SEQ ID NO:45), EIQSLV (SEQ ID NO:46), NEIQSLV (SEQ ID NO:47), and RNEIQSLV (SEQ ID NO:48) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD97 is ESGI (SEQ ID NO:49). When naturally-occurring residues are added or removed from the core sequence, GI (SEQ ID NO:), SGI (SEQ ID NO:), SESGI (SEQ ID NO:50), ASESGI (SEQ ID NO:51), RASESGI (SEQ ID NO:52), and LRASESGI (SEQ ID NO:53) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD98 is PYAA (SEQ ID NO:54). When naturally-occurring residues are added or removed from the core sequence, AA (SEQ ID NO:), YAA (SEQ ID NO:), FPYAA (SEQ ID NO:55), RFPYAA (SEQ ID NO:56), LRFPYAA (SEQ ID NO:57), and LLRFPYAA (SEQ ID NO:58) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CDw137 is GCEL (SEQ ID NO:59). When naturally-occurring residues are added or removed from the core sequence, EL (SEQ ID NO:), CEL (SEQ ID NO:), GGCEL (SEQ ID NO:60), EGGCEL (SEQ ID NO:61), EEGGCEL (SEQ ID NO:62), and EEGGCEL (SEQ ID NO:63) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CD166 is KTEA (SEQ ID NO:64). When naturally-occurring residues are added or removed from the core sequence, EA (SEQ ID NO:), TEA (SEQ ID NO:), HKTEA (SEQ ID NO:65), NHKTEA (SEQ ID NO:66), NNHKTEA (SEQ ID NO:67), and ENNHKTEA (SEQ ID NO:68) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CDw128 is SSNL (SEQ ID NO:69). When naturally-occurring residues are added or removed from the core sequence, NL (SEQ ID NO:), SNL (SEQ ID NO:), VSSNL (SEQ ID NO:70), NVSSNL (SEQ ID NO:71), VNVSSNL (SEQ ID NO:72), and SVNSSNL (SEQ ID NO:73) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of DNAM-1 is KTRV (SEQ ID NO:74). When naturally-occurring residues are added or removed from the core sequence, RV (SEQ ID NO:), TRV (SEQ ID NO:), PKTRV (SEQ ID NO:75), RPKTRV (SEQ ID NO:76), RRPKTRV (SEQ ID NO:77), and SRRPKTRV (SEQ ID NO:78) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of FasL is LYKL (SEQ ID NO:79). When naturally-occurring residues are added or removed from the core sequence, KL (SEQ ID NO:), YKL (SEQ ID NO:), GLYKL (SEQ ID NO:80), FGLYKL (SEQ ID NO:81), FFGLYKL (SEQ ID NO:82), and TFFGLYKL (SEQ ID NO:83) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of LPAP is VTAL (SEQ ID NO:84). When naturally-occurring residues are added or removed from the core sequence, AL (SEQ ID NO:), TAL (SEQ ID NO:), HVTAL (SEQ ID NO:85), LHVTAL (SEQ ID NO:86), GLHVTAL (SEQ ID NO:87), and QGLHVTAL (SEQ ID NO:88) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CLASP-1 is SAQV (SEQ. ID. NO: 182). When naturally-occurring residues are added or removed from the core sequence, QV (SEQ ID NO:), AQV (SEQ ID NO:), SSAQV (SEQ. ID. NO: 183), SSSAQV (SEQ. ID. NO: 184), ISSSAQV (SEQ. ID. NO: 185), and SISSSAQV (SEQ. ID. NO: 186) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CLASP-2 is SSVV (SEQ. ID. NO: 187). When naturally-occurring residues are added or removed from the core sequence, VV (SEQ ID NO:), SVV (SEQ ID NO:), SSSVV (SEQ. ID. NO: 188), SSSSVV (SEQ. ID. NO: 189), TSSSVV (SEQ. ID. NO: 190), and MTSSSVV (SEQ. ID. NO: 191) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of CLASP-4 is YAEV (SEQ. ID. NO: 192). When naturally-occurring residues are added or removed from the core sequence, EV (SEQ ID NO:), AEV (SEQ ID NO:), RYAEV (SEQ. ID. NO: 193), PRYAEV (SEQ. ID. NO: 194), SPRYAEV (SEQ. ID. NO: 195), and GSPRYAEV (SEQ. ID. NO: 196) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of KV1.3 is FTDV (SEQ. ID. NO: 202). When naturally-occurring residues are added or removed from the core sequence, DV (SEQ ID NO:), TDV (SEQ ID NO:), IFTDV (SEQ. ID. NO: 203), KIFTDV (SEQ. ID. NO: 204), KKIFTDV (SEQ. ID. NO: 205), and IKKIFTDV (SEQ. ID. NO: 206) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of DOCK2 is STD L (SEQ. ID. NO: 207). When naturally-occurring residues are added or removed from the core sequence, DL (SEQ. ID. NO:), TDL (SEQ. ID. NO:), LSTD L (SEQ. ID. NO: 208), SLSTD L (SEQ. ID. NO: 209), DSLSTD L (SEQ. ID. NO: 210), and PDSLSTD L (SEQ. ID. NO: 211) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of BLR-1 is LTTF (SEQ ID NO:). When naturally-occurring residues are added or removed from the core sequence, TF (SEQ ID NO:), TTF (SEQ ID NO:), SLTTF (SEQ ID NO:), TSLTTF (SEQ ID NO:), ATSLTTF (SEQ ID NO:), and NATSLTTF (SEQ ID NO:) may also be used to target a PDZ domain-containing protein in T cells.

The C-terminal core sequence of PAG is ITRL (SEQ ID NO:). When naturally-occurring residues are added or removed from the core sequence, RL (SEQ ID NO:), TRL (SEQ ID NO:), DITRL (SEQ ID NO:), RDITRL (SEQ ID NO:), GRDITRL (SEQ ID NO:), and QGRDITRL (SEQ ID NO:) may also be used to target a PDZ domain-containing protein in T cells.

6.6.2 PDZ Motif Sequences of B Cell Surface Receptors

A number of surface receptors expressed by B cells contain a PDZ domain motif sequence. These molecules include, but are not limited to, CD38, CD53, CD95, CD97, CD98, CDw137, CD138, CDw125 (IL5R), DNAM-1, LPAP, Syndecan-2 (Barclay et al., 1997, The Leucocyte Antigen Facts Book, second edition, Academic Press) and BLR-1. The specific motif sequences of CD38, CD53, CD83, CD95, CD97, CD98, CDw137, DNAM-1, DOCK2, LPAP, BLR-1 (CXCR5), PAG, CLASP-1, CLASP-2 and CLASP-4 have been described in the preceding paragraphs.

The C-terminal core sequence of CD138 is EFYA (SEQ ID NO:89). When naturally-occurring residues are added or removed from the core sequence, YA (SEQ ID NO:), FYA (SEQ ID NO:), EEFYA (SEQ ID NO:90), QEEFYA (SEQ ID NO:91), KQEEFYA (SEQ ID NO:92), and TKQEEFYA (SEQ ID NO:93) may also be used to target a PDZ domain-containing protein in B cells.

The C-terminal core sequence of CDw125 is DSVF (SEQ ID NO:94). When naturally-occurring residues are added or removed from the core sequence, VF (SEQ ID NO:), SVF (SEQ ID NO:), EDSVF (SEQ ID NO:95), LEDSVF (SEQ ID NO:96), TLEDSVF (SEQ ID NO:97), and ETLEDSVF (SEQ ID NO:98) may also be used to target a PDZ domain-containing protein in B cells.

The C-terminal core sequence of Syndecan-2 is EFYA (SEQ. ID. NO: 212). When naturally-occurring residues are added or removed from the core sequence, YA (SEQ ID NO:), FYA (SEQ ID NO:), KEFYA (SEQ. ID. NO: 213), TKEFYA (SEQ. ID. NO: 214), PTKEFYA (SEQ. ID. NO: 215), and APTKEFYA (SEQ. ID. NO: 216) may also be used to target a PDZ domain-containing protein in B cells.

The C-terminal core sequence of BLR-1 is LTTF (SEQ. ID. NO: 217). When

naturally-occurring residues are added or removed from the core sequence, TF (SEQ ID NO:), TTF (SEQ ID NO:), SLTTF (SEQ. ID. NO: 218), TSLTTF (SEQ. ID. NO: 219), ATSLTTF (SEQ. ID. NO: 220), and NATSLTTF (SEQ. ID. NO: 221) may also be used to target a PDZ domain-containing protein in B cells.

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6.6.3 PDZ motif sequences of Natural Killer Cell Surface Receptors

A number of surface receptors expressed by NK cells contain a PDZ domain motif sequence. These molecules include, but are not limited to CD38, CD56, CD98 and DNAM-1. The specific motif sequences of CD38, CD98 and DNAM-1 have been described

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in the preceding paragraphs. The C-terminal core sequence of CD56 is ESKA (SEQ ID NO:99). When naturally-occurring residues are added or removed from the core sequence, KA (SEQ ID NO:), SKA (SEQ ID NO:), NESKA (SEQ ID NO:100), ENESKA (SEQ ID NO:101), KENESKA (SEQ ID NO:102), and TKENESKA (SEQ ID NO:103) may also be used to target a PDZ

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6.6.4 PDZ motif sequences of Monocyte Surface Receptors

A number of surface receptors expressed by cells of the monocytic lineage (monocytes and macrophages) contain a PDZ domain motif sequence. These molecules include, but are not limited to CD38, CD44, CD46, CD49e, CD49f, CD53, CD61, CD95, CD97, CD98, CD148, CDw128, CDw137, Ly-6, DNAM-1 and FcεRIβ. The specific motif sequences of CD38, CD49e, CD49f, CD53, CD95, CD97, CD98, CDw128, CDw137, DNAM-1, Galectin 3 (Mac-2), BLR-1 (CXCR5) and Mannose receptor have been described in the preceding paragraphs.

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The C-terminal core sequence of CD44 is KIGV (SEQ ID NO:104). When naturally-occurring residues are added or removed from the core sequence, GV (SEQ ID NO:), IGV (SEQ ID NO:), MKIGV (SEQ ID NO:105), DMKIGV (SEQ ID NO:106), VDMKIGV (SEQ ID NO:107) and NVDMKIGV (SEQ ID NO:108) may also be used to target a PDZ domain-containing protein in monocytes.

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The C-terminal core sequence of CD46 is FTSL (SEQ ID NO:109). When naturally-occurring residues are added or removed from the core sequence, SL (SEQ ID NO:), TSL (SEQ ID NO:), KFTSL (SEQ ID NO:110), VKFTSL (SEQ ID NO:111), EVKFTSL (SEQ ID NO:112) and REVKFTSL (SEQ ID NO:113) may also be used to target a PDZ domain-containing protein in monocytes.

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The C-terminal core sequence of CD61 is KSLV (SEQ ID NO:114). When

naturally-occurring residues are added or removed from the core sequence, LV (SEQ ID NO:), SLV (SEQ ID NO:), LKSLV (SEQ ID NO:115), FLKSLV (SEQ ID NO:116), RFLKSLV (SEQ ID NO:117) and GRFLKSLV (SEQ ID NO:118) may also be used to target a PDZ domain-containing protein in monocytes.

5 The C-terminal core sequence of CD148 is GYIA (SEQ ID NO:119). When naturally-occurring residues are added or removed from the core sequence, IA (SEQ ID NO:), YIA (SEQ ID NO:), NGYIA (SEQ ID NO:120), TNGYIA (SEQ ID NO:121), KTNKYIA (SEQ ID NO:122) and GKTNGYIA (SEQ ID NO:123) may also be used to target a PDZ domain-containing protein in monocytes.

10 The C-terminal core sequence of Ly-6 is QTLL (SEQ ID NO:124). When naturally-occurring residues are added or removed from the core sequence, LL (SEQ ID NO:), TLL (SEQ ID NO:), LQTLL (SEQ ID NO:125), LLQTLL (SEQ ID NO:126), VLLQTLL (SEQ ID NO:127) and SVLLQTLL (SEQ ID NO:128) may also be used to target a PDZ domain-containing protein in monocytes.

15 The C-terminal core sequence of FcεRIβ is PIDL (SEQ ID NO:129). When naturally-occurring residues are added or removed from the core sequence, DL (SEQ ID NO:), IDL (SEQ ID NO:), PPIDL (SEQ ID NO:130), SPPIDL (SEQ ID NO:131), MSPPIDL (SEQ ID NO:132) and EMSPPIDL (SEQ ID NO:133) may also be used to target a PDZ domain-containing protein in monocytes.

20 The C-terminal core sequence of Galectin 3 is YTMI (SEQ ID NO:134). When naturally-occurring residues are added or removed from the core sequence, MI (SEQ ID NO:), TMI (SEQ ID NO:), SYTMI (SEQ ID NO:135), ASYTMI (SEQ ID NO:136), SASYTMI (SEQ ID NO:137) and TSASYTMI (SEQ ID NO:138) may also be used to target a PDZ domain-containing protein in monocytes.

25 The C-terminal core sequence of mannose receptor is HSVI (SEQ ID NO:139). When naturally-occurring residues are added or removed from the core sequence, VI (SEQ ID NO:), SVI (SEQ ID NO:), EHSVI (SEQ ID NO:140), NEHSVI (SEQ ID NO:141), QNEHSVI (SEQ ID NO:142) and EQNEHSVI (SEQ ID NO:143) may also be used to target a PDZ domain-containing protein in monocytes.

30 6.6.5 PDZ motif sequences of Granulocyte Surface Receptors

A number of surface receptors expressed by granulocytes contain a PDZ domain motif sequence. These molecules include, but are not limited to CD53, CD95, CD97, CD98, CD148, CDw125, CDw128, FcεRIβ and G-CSFR. The specific motif sequences of most of these molecules have been described in the preceding paragraphs.

The C-terminal core sequence of G-CSFR is TSVL (SEQ ID NO:144). When naturally-occurring residues are added or removed from the core sequence, VL (SEQ ID NO:), SVL (SEQ ID NO:), ITSVL (SEQ ID NO:145), PITSVL (SEQ ID NO:146), FPITSVL (SEQ ID NO:147) and LFPITSVL (SEQ ID NO:148) may also be used to target a PDZ domain-containing protein in monocytes.

6.6.6 PDZ motif sequences of Endothelial Cell Surface Receptors

While endothelial cells are not hematopoietic cells, they closely interact with the hematopoietic system as they form the lining of blood vessels. As such, endothelial cells come in contact with the cells of the hematopoietic system. Thus, the ability to regulate endothelial cell function provides for indirect regulation of hematopoietic cells.

A number of surface receptors expressed by endothelial cells contain a PDZ domain motif sequence. These molecules include, but are not limited to CD34, CD46, CD66b, CD66c, CD105, CD106, CD62e (E-selectin) and VCAM1.

The C-terminal core sequence of CD34 is DTEL (SEQ ID NO:149). When naturally-occurring residues are added or removed from the core sequence, EL (SEQ ID NO:), TEL (SEQ ID NO:), ADTEL (SEQ ID NO:150), VADTEL (SEQ ID NO:151), VVADTEL (SEQ ID NO:152) and HVVADTEL (SEQ ID NO:153) may also be used to target a PDZ domain-containing protein in endothelial cells.

The C-terminal core sequence of CD66b and CD66c is VALI (SEQ ID NO:154). When naturally-occurring residues are added or removed from the core sequence, LI (SEQ ID NO:), ALI (SEQ ID NO:), RVALI (SEQ ID NO:155), ARVALI (SEQ ID NO:156), LARVALI (SEQ ID NO:157) and VLARVALI (SEQ ID NO:158) may also be used to target a PDZ domain-containing protein in endothelial cells.

The C-terminal core sequence of CD105 is SSMA (SEQ ID NO:159). When naturally-occurring residues are added or removed from the core sequence, MA (SEQ ID NO:), SMA (SEQ ID NO:), TSSMA (SEQ ID NO:160), STSSMA (SEQ ID NO:161), CSTSSMA (SEQ ID NO: 222) and PCSTSSMA (SEQ ID NO: 162) may also be used to target a PDZ domain-containing protein in endothelial cells.

The C-terminal core sequence of CD106 is KSKV (SEQ ID NO:163). When naturally-occurring residues are added or removed from the core sequence, KV (SEQ ID NO:), SKV (SEQ ID NO:), QKSKV (SEQ ID NO:164), AQKSKV (SEQ ID NO:165), EAQKSKV (SEQ ID NO:166) and VEAQKSKV (SEQ ID NO:167) may also be used to target a PDZ domain-containing protein in endothelial cells.

The C-terminal core sequence of CD62e is SYIL (SEQ ID NO:168). When naturally-occurring residues are added or removed from the core sequence, IL (SEQ ID NO:), YIL (SEQ ID NO:), PSYIL (SEQ ID NO:169), KPSYIL (SEQ ID NO:170), QKPSYIL (SEQ ID NO:171) and YQKPSYIL (SEQ ID NO:172) may also be used to target a PDZ domain-containing protein in endothelial cells.

The C-terminal core sequence of VCAM1 is KSKV (SEQ. ID. NO: 197). When naturally-occurring residues are added or removed from the core sequence, KV (SEQ ID NO:), SKV (SEQ ID NO:), QKSKV (SEQ. ID. NO: 198), AQKSKV (SEQ. ID. NO: 199), EAQKSKV (SEQ. ID. NO: 200), and VEAQKSKV (SEQ. ID. NO: 201) may also be used to target a PDZ domain-containing protein in endothelial cells.

6.6.7 Mast Cell, Basophils and Eosinophil Cell Surface Receptors

FcεRIβ, CDw125, CDw128 and IL-8RB are transmembrane receptors expressed by mast cells, basophils and eosinophils. These receptors play a role in the activation of these cells to result in degranulation and histamine release in allergic reactions. The C-terminal core sequence of FcεRIβ is PIDL (SEQ ID NO:129). When naturally-occurring residues are added or removed from the core sequence, DL (SEQ ID NO:), IDL (SEQ ID NO:), PPIDL (SEQ ID NO:244), SPPIDL (SEQ ID NO:245), MSPPIDL (SEQ ID NO:246) and EMSPPIDL (SEQ ID NO:247) may also be used to target a PDZ domain-containing protein in mast cells. In addition, the residue E may be substituted with G to increase its binding affinity.

The C-terminal core sequence of CDw125 is DSVF (SEQ ID NO: 248). When naturally-occurring residues are added or removed from the core sequence, VF (SEQ ID NO:), SVF (SEQ ID NO:), EDSVF (SEQ ID NO:249), LEDSVF (SEQ ID NO:250), TLEDVF (SEQ ID NO:251), and ETLEDVF (SEQ ID NO:252) may also be used to target a PDZ domain-containing protein in mast cells.

The C-terminal core sequence of CDw128 is SSNL (SEQ ID NO:253). When naturally-occurring residues are added or removed from the core sequence, NL (SEQ ID NO:), SNL (SEQ ID NO:), VSSNL (SEQ ID NO:254), NVSSNL (SEQ ID NO:255), VNVSSNL (SEQ ID NO:256), and SVNVSNNL (SEQ ID NO:257) may also be used to target a PDZ domain-containing protein in mast cells.

The C-terminal core sequence of IL-8RB is STTL (SEQ ID NO:258). When naturally-occurring residues are added or removed from the core sequence, TL (SEQ ID NO:), TTL (SEQ ID NO:), TSTTL (SEQ ID NO:259), HTSTTL (SEQ ID NO:260), GHTSTTL (SEQ ID NO:261) and SGHTSTTL (SEQ ID NO:262) may also be used to target a PDZ domain-containing protein in mast cells.

6.6.8 Other PDZ motif sequences

The C-terminal core sequence of NMDA is ESDV (SEQ. ID. NO: 223). When naturally-occurring residues are added or removed from the core sequence, DV (SEQ ID NO:), SDV (SEQ ID NO:), IESDV (SEQ. ID. NO: 224), SIESDV (SEQ. ID. NO: 225), PSIESDV (SEQ. ID. NO: 226), and MPSIESDV (SEQ. ID. NO: 227) may also be used to target a PDZ domain-containing protein in neuronal cells.

The C-terminal core sequence of neurexin is EYYV (SEQ. ID. NO: 228). When naturally-occurring residues are added or removed from the core sequence, YV (SEQ ID NO:), YYV (SEQ ID NO:), KEYV (SEQ. ID. NO: 229), DKEYV (SEQ. ID. NO: 230), KDKEYV (SEQ. ID. NO: 231), and NKDKEYV (SEQ. ID. NO: 232) may also be used to target a PDZ domain-containing protein in neuronal cells.

The C-terminal core sequence of Glycophorin C is EYFI (SEQ. ID. NO: 233). When naturally-occurring residues are added or removed from the core sequence, FI (SEQ ID NO:), YFI (SEQ ID NO:), KEYFI (SEQ. ID. NO: 234), RKEYFI (SEQ. ID. NO: 235), SRKEYFI (SEQ. ID. NO: 236), and SSRKEYFI (SEQ. ID. NO: 237) may also be used to target a PDZ domain-containing protein.

The C-terminal core sequence of CD148 is KTIA (SEQ. ID. NO: 238). When naturally-occurring residues are added or removed from the core sequence, IA (SEQ ID NO:), TIA (SEQ ID NO:), GKTIA (SEQ. ID. NO: 239), FGKTIA (SEQ. ID. NO: 240), TFGKTIA (SEQ. ID. NO: 241), and TTFGKTIA (SEQ. ID. NO: 242) may also be used to target a PDZ domain-containing protein in epithelial or myeloid cells.

The C-terminal core sequence of beta-spectrin is VSFV (SEQ. ID. NO:). When naturally-occurring residues are added to the core sequence, FV (SEQ. ID. NO:), SFV (SEQ. ID. NO:), LVSFV (SEQ. ID. NO:), SLVSFV (SEQ. ID. NO:), QSLVSFV (SEQ. ID. NO:) AND GQSLVSFV (SEQ. ID. NO:) may also be used to target a PDZ domain-containing protein.

6.7. Preparation of Peptides

6.7.1. Chemical Synthesis

The peptides of the invention or analogues thereof, may be prepared using virtually any art-known technique for the preparation of peptides and peptide analogues. For example, the peptides may be prepared in linear form using conventional solution or solid phase

peptide syntheses and cleaved from the resin followed by purification procedures (Creighton, 1983, Protein Structures And Molecular Principles, W.H. Freeman and Co., N.Y.). Suitable procedures for synthesizing the peptides described herein are well known in the art. The composition of the synthetic peptides may be confirmed by amino acid analysis or sequencing (e.g., the Edman degradation procedure and mass spectroscopy).

In addition, analogues and derivatives of the peptides can be chemically synthesized. The linkage between each amino acid of the peptides of the invention may be an amide, a substituted amide or an isostere of amide. Nonclassical amino acids or chemical amino acid analogues can be introduced as a substitution or addition into the sequence. Non-classical amino acids include, but are not limited to, the D-isomers of the common amino acids, α -amino isobutyric acid, 4-aminobutyric acid, Abu, 2-amino butyric acid, γ -Abu, ϵ -Ahx, 6-amino hexanoic acid, Aib, 2-amino isobutyric acid, 3-amino propionic acid, ornithine, norleucine, norvaline, hydroxyproline, sarcosine, citrulline, cysteic acid, t-butylglycine, t-butylalanine, phenylglycine, cyclohexylalanine, β -alanine, fluoro-amino acids, designer amino acids such as β -methyl amino acids, $C\alpha$ -methyl amino acids, $N\alpha$ -methyl amino acids, and amino acid analogues in general. Furthermore, the amino acid can be D (dextrorotary) or L (levorotary).

6.7.2. Recombinant Synthesis

If the peptide is composed entirely of gene-encoded amino acids, or a portion of it is so composed, the peptide or the relevant portion may also be synthesized using conventional recombinant genetic engineering techniques. For recombinant production, a polynucleotide sequence encoding a linear form of the peptide is inserted into an appropriate expression vehicle, *i.e.*, a vector which contains the necessary elements for the transcription and translation of the inserted coding sequence, or in the case of an RNA viral vector, the necessary elements for replication and translation. The expression vehicle is then transfected into a suitable target cell which will express the peptide. Depending on the expression system used, the expressed peptide is then isolated by procedures well-established in the art. Methods for recombinant protein and peptide production are well known in the art (*see, e.g.*, Maniatis *et al.*, 1989, Molecular Cloning A Laboratory Manual, Cold Spring Harbor Laboratory, N.Y.; and Ausubel *et al.*, 1989, Current Protocols in Molecular Biology, Greene Publishing Associates and Wiley Interscience, N.Y.).

5 A variety of host-expression vector systems may be utilized to express the peptides described herein. These include, but are not limited to, microorganisms such as bacteria transformed with recombinant bacteriophage DNA or plasmid DNA expression vectors containing an appropriate coding sequence; yeast or filamentous fungi transformed with recombinant yeast or fungi expression vectors containing an appropriate coding sequence; insect cell systems infected with recombinant virus expression vectors (*e.g.*, baculovirus) containing an appropriate coding sequence; plant cell systems infected with recombinant virus expression vectors (*e.g.*, cauliflower mosaic virus or tobacco mosaic virus) or transformed with recombinant plasmid expression vectors (*e.g.*, Ti plasmid) containing an appropriate coding sequence; or animal cell systems.

10 The expression elements of the expression systems vary in their strength and specificities. Depending on the host/vector system utilized, any of a number of suitable transcription and translation elements, including constitutive and inducible promoters, may be used in the expression vector. For example, when cloning in bacterial systems, inducible promoters such as pL of bacteriophage λ , plac, ptrp, ptac (ptrp-lac hybrid promoter) and the like
15 may be used; when cloning in insect cell systems, promoters such as the baculovirus polyhedron promoter may be used; when cloning in plant cell systems, promoters derived from the genome of plant cells (*e.g.*, heat shock promoters; the promoter for the small subunit of RUBISCO; the promoter for the chlorophyll a/b binding protein) or from plant viruses (*e.g.*, the 35S RNA promoter of CaMV; the coat protein promoter of TMV) may be used; when
20 cloning in mammalian cell systems, promoters derived from the genome of mammalian cells (*e.g.*, metallothionein promoter) or from mammalian viruses (*e.g.*, the adenovirus late promoter; the vaccinia virus 7.5 K promoter) may be used; when generating cell lines that contain multiple copies of expression product, SV40-, BPV- and EBV-based vectors may be used with
25 an appropriate selectable marker.

In cases where plant expression vectors are used, the expression of sequences encoding the peptides of the invention may be driven by any of a number of promoters. For example, viral promoters such as the 35S RNA and 19S RNA promoters of CaMV (Brisson *et al.*, 1984, Nature 310:511-514), or the coat protein promoter of TMV (Takamatsu *et al.*, 1987, EMBO J. 6:307-311) may be used; alternatively, plant promoters such as the small subunit of
30 RUBISCO (Coruzzi *et al.*, 1984, EMBO J. 3:1671-1680; Broglie *et al.*, 1984, Science 224:838-843) or heat shock promoters, *e.g.*, soybean hsp17.5-E or hsp17.3-B (Gurley *et al.*, 1986, Mol.

Cell. Biol. 6:559-565) may be used. These constructs can be introduced into planleukocytes using Ti plasmids, Ri plasmids, plant virus vectors, direct DNA transformation, microinjection, electroporation, etc. For reviews of such techniques *see, e.g.*, Weissbach & Weissbach, 1988, Methods for Plant Molecular Biology, Academic Press, NY, Section VIII, pp. 421-463; and
5 Grierson & Corey, 1988, Plant Molecular Biology, 2d Ed., Blackie, London, Ch. 7-9.

In one insect expression system that may be used to produce the peptides of the invention, *Autographa californica* nuclear polyhidrosis virus (AcNPV) is used as a vector to express the foreign genes. The virus grows in *Spodoptera frugiperda* cells. A coding sequence may be cloned into non-essential regions (for example the polyhedron gene) of the virus and
10 placed under control of an AcNPV promoter (for example, the polyhedron promoter). Successful insertion of a coding sequence will result in inactivation of the polyhedron gene and production of non-occluded recombinant virus (*i.e.*, virus lacking the proteinaceous coat coded for by the polyhedron gene). These recombinant viruses are then used to infect *Spodoptera frugiperda* cells in which the inserted gene is expressed. (*e.g.*, see Smith *et al.*, 1983, J. Virol.
15 46:584; Smith, U.S. Patent No. 4,215,051). Further examples of this expression system may be found in Current Protocols in Molecular Biology, Vol. 2, Ausubel *et al.*, eds., Greene Publish. Assoc. & Wiley Interscience.

In mammalian host cells, a number of viral based expression systems may be utilized. In cases where an adenovirus is used as an expression vector, a coding sequence may
20 be ligated to an adenovirus transcription/translation control complex, *e.g.*, the late promoter and tripartite leader sequence. This chimeric gene may then be inserted in the adenovirus genome by *in vitro* or *in vivo* recombination. Insertion in a non-essential region of the viral genome (*e.g.*, region E1 or E3) will result in a recombinant virus that is viable and capable of expressing peptide in infected hosts. (*e.g.*, See Logan & Shenk, 1984, Proc. Natl. Acad. Sci. USA
25 81:3655-3659). Alternatively, the vaccinia 7.5 K promoter may be used, (*see, e.g.*, Mackett *et al.*, 1982, Proc. Natl. Acad. Sci. USA 79:7415-7419; Mackett *et al.*, 1984, J. Virol. 49:857-864; Panicali *et al.*, 1982, Proc. Natl. Acad. Sci. USA 79:4927-4931).

Other expression systems for producing linear peptides of the invention will be apparent to those having skill in the art.

30 6.7.3. Purification of the Peptides and Peptide Analogues

The peptides and peptide analogues of the invention can be purified by art-known techniques such as high performance liquid chromatography, ion exchange chromatography, gel electrophoresis, affinity chromatography and the like. The actual conditions used to purify a particular peptide or analogue will depend, in part, on factors such as net charge, hydrophobicity, hydrophilicity, etc., and will be apparent to those having skill in the art. The purified peptides can be identified by assays based on their physical or functional properties, including radioactive labeling followed by gel electrophoresis, radioimmuno-assays, ELISA, bioassays, and the like.

For affinity chromatography purification, any antibody which specifically binds the peptides or peptide analogues may be used. For the production of antibodies, various host animals, including but not limited to rabbits, mice, rats, etc., may be immunized by injection with a peptide. The peptide may be attached to a suitable carrier, such as BSA or KLH, by means of a side chain functional group or linkers attached to a side chain functional group. Various adjuvants may be used to increase the immunological response, depending on the host species, including but not limited to Freund's (complete and incomplete), mineral gels such as aluminum hydroxide, surface active substances such as lysolecithin, pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, dinitrophenol, and potentially useful human adjuvants such as BCG (bacilli Calmette-Guerin) and *Corynebacterium parvum*.

Monoclonal antibodies to a peptide may be prepared using any technique which provides for the production of antibody molecules by continuous cell lines in culture. These include but are not limited to the hybridoma technique originally described by Koehler and Milstein, 1975, Nature 256:495-497, the human B-cell hybridoma technique, Kosbor *et al.*, 1983, Immunology Today 4:72; Cote *et al.*, 1983, Proc. Natl. Acad. Sci. U.S.A. 80:2026-2030 and the EBV-hybridoma technique (Cole *et al.*, 1985, Monoclonal Antibodies and Cancer Therapy, Alan R. Liss, Inc., pp. 77-96 (1985)). In addition, techniques developed for the production of "chimeric antibodies" (Morrison *et al.*, 1984, Proc. Natl. Acad. Sci. U.S.A. 81:6851-6855; Neuberger *et al.*, 1984, Nature 312:604-608; Takeda *et al.*, 1985, Nature 314:452-454) by splicing the genes from a mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity can be used. Alternatively, techniques described for the production of single chain antibodies (U.S. Patent No. 4,946,778) can be adapted to produce peptide-specific single chain antibodies.

Antibody fragments which contain deletions of specific binding sites may be generated by known techniques. For example, such fragments include but are not limited to F(ab')₂ fragments, which can be produced by pepsin digestion of the antibody molecule and Fab fragments, which can be generated by reducing the disulfide bridges of the F(ab')₂ fragments.

5 Alternatively, Fab expression libraries may be constructed (Huse *et al.*, 1989, Science 246:1275-1281) to allow rapid and easy identification of monoclonal Fab fragments with the desired specificity for the peptide of interest.

The antibody or antibody fragment specific for the desired peptide can be attached, for example, to agarose, and the antibody-agarose complex is used in
10 immunochromatography to purify peptides of the invention. See, Scopes, 1984, Protein Purification: Principles and Practice, Springer-Verlag New York, Inc., NY, Livingstone, 1974, Methods Enzymology: Immunoaffinity Chromatography of Proteins 34:723-731.

6.8. Uses of PDZ Domain Binding and Antagonist Compounds

In one aspect of the invention, the PDZ domain binding and PDZ-PL inhibitory
15 compounds of the present invention are useful in regulating diverse activities of hematopoietic cells (e.g., T cells and B cells) and other cells involved in the immune response.

In one embodiment of the invention, the compounds of the invention are used to inhibit leukocyte activation, which is manifested in measurable events including but not limited to, cytokine production, cell adhesion, expansion of cell numbers, apoptosis and
20 cytotoxicity. As a corollary, the compounds of the invention may be used to treat diverse conditions associated with undesirable leukocyte activation, including but not limited to, acute and chronic inflammation, graft-versus-host disease, transplantation rejection, hypersensitivities and autoimmunity such as multiple sclerosis, rheumatoid arthritis, periodontal disease, systemic lupus erythematosus, juvenile diabetes mellitus, non-insulin-dependent diabetes, and allergies,
25 and other conditions listed herein (see, e.g., Section 6.4, *supra*).

Thus, the invention also relates to methods of using such compositions in modulating leukocyte activation as measured by, for example, cytotoxicity, cytokine production, cell proliferation, and apoptosis. Assays for activation are well known. For example, PDZ/PL interaction antagonists can be evaluated in the following: (1) cytotoxic T
30 lymphocytes can be incubated with radioactively labeled target cells and the antigen-specific lysis of these target cells detected by the release of radioactivity, (2) helper T lymphocytes can

be incubated with antigens and antigen presenting cells and the synthesis and secretion of cytokines measured by standard methods (Windhagen A; et al., 1995, Immunity 2(4): 373-80), (3) antigen presenting cells can be incubated with whole protein antigen and the presentation of that antigen on MHC detected by either T lymphocyte activation assays or biophysical methods (Harding et al., 1989, Proc. Natl. Acad. Sci., 86: 4230-4), (4) mast cells can be incubated with reagents that cross-link their Fc-epsilon receptors and histamine release measured by enzyme immunoassay (Siraganian, et al., 1983, TIPS 4: 432-437).

Similarly, the effect of PDZ/PL interaction antagonists on products of leukocyte activation in either a model organism (e.g., mouse) or a human patient can also be evaluated by various methods that are well known. For example, (1) the production of antibodies in response to vaccination can be readily detected by standard methods currently used in clinical laboratories, e.g., an ELISA; (2) the migration of immune cells to sites of inflammation can be detected by scratching the surface of skin and placing a sterile container to capture the migrating cells over scratch site (Peters et al., 1988, Blood 72: 1310-5); (3) the proliferation of peripheral blood mononuclear cells in response to mitogens or mixed lymphocyte reaction can be measured using ³H-thymidine; (4) the phagocytic capacity of granulocytes, macrophages, and other phagocytes in PBMCs can be measured by placing PMBCs in wells together with labeled particles (Peters et al., 1988); and (5) the differentiation of immune system cells can be measured by labeling PBMCs with antibodies to CD molecules such as CD4 and CD8 and measuring the fraction of the PBMCs expressing these markers.

In one exemplary assay, human peripheral blood mononuclear cells (PBMC), human T cell clones (e.g., Jurkat E6, ATCC TIB-152), EBV-transformed B cell clones (e.g., 9D10, ATCC CRL-8752), antigen-specific T cell clones or lines can be used to examine PDZ/PL interaction antagonists in vitro. Inhibition of activation of these cells or cell lines can be used for the evaluation of potential PDZ/PL interaction antagonists.

Standard methods by which hematopoietic cells are stimulated to undergo activation characteristic of an immune response are, for example:

A) Antigen specific stimulation of immune responses. Either pre-immunized or naïve mouse splenocytes can be generated by standard procedures. In addition, antigen-specific T cell clones and hybridomas (e.g., MBP-specific), and numerous B cell lymphoma cell lines (e.g., CH27), have been previously characterized and are available for the assays discussed below. Antigen specific splenocytes or B-cells can be mixed with antigen specific T-cells in

responses are generated in the presence of potential PDZ/PL interaction antagonists is indicative of a potential PDZ/PL interaction antagonists.

(B) Intracellular Calcium Flux

The kinetics of intracellular Ca^{2+} concentrations are measured over time after stimulation of cells preloaded with a calcium sensitive dye. Upon binding Ca^{2+} the indicator dye (e.g., Fluor-4 (Molecular Probes)), exhibits an increase in fluorescence level using flow cytometry, solution fluorometry, and confocal microscopy. Any change in the level or timing of calcium flux when immune responses are generated in the presence of PDZ/PL interaction antagonists is indicative of an inhibition of this response.

10 (C) Regulation of early activation markers

Increased and diminished expression/regulation of early lymphocyte activation marker levels such as CD69, IL-2R, MHC class II, B7, and TCR are commonly measured with fluorescently labeled antibodies using flow cytometry. All antibodies are commercially available. Any change in the expression levels of lymphocyte activation markers when immune responses are generated in the presence of the PDZ/PL interaction antagonists is indicative of an inhibition of this response.

(D) Increased metabolic activity/acid release

Activation of most known signal transduction pathways trigger increases in acidic metabolites. This reproducible biological event is measured as the rate of acid release using a microphysiometer (Molecular Devices), and is used as an early activation marker when comparing the treatment of cells with potential biological therapeutics (McConnell, H.M. et al., 1992, *Science* 257: 1906-1912 and McConnell, H.M., 1995, *Proc. Natl. Acad. Sci.* 92: 2750-2754). Any statistically significant increase or decrease in acid release of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function.

(E) Cell proliferation/cell viability assays

(1) ^3H -thymidine incorporation

Exposure of lymphocytes to antigen or mitogen in vitro induces DNA synthesis and cellular proliferation. The measurement of mitotic activity by ^3H -thymidine incorporation into newly synthesized DNA is one of the most frequently used assays to quantitative T cell activation. Depending on the cell population and form of stimulation used to activate the T cells, mitotic activity can be measured within 24-72 hrs. in vitro, post ^3H -thymidine pulse



(Mishell, B. B. and S. M. Shiigi, 1980, Selected Methods in Cellular Immunology, W. H. Freeman and Company and Dutton, R. W. and Pearce, J. D., 1962, Nature 194: 93). Any statistically significant increase or decrease in CPM of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function.

(2) MTS [5-(3-carboxymethoxyphenyl)-2-(4,5-dimethylthiazolyl)-3(4-sulfophenyl)tetrazolium, inner salt] is a colorimetric method for determining the number of viable cells in proliferation or cytotoxicity assays (Barltrop, J.A. et al., 1991, Bioorg. & Med. Chem. Lett. 1: 611). 1-5 days after lymphocyte activation, MTS tetrazolium compound, Owen's reagent, is bio-reduced by cells into a colored formazan product that is soluble in tissue culture media. Color intensity is read at 490 nm minus 650 nm using a microplate reader. Any statistically significant increase or decrease in color intensity of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), can suggest an effect of the PDZ/PL interaction antagonist on biological function (Mosmann, T., 1983, J. Immunol. Methods 65: 55 and Barltrop, J.A. et al. (1991)).

(3) Bromodeoxyuridine (BrdU), a thymidine analogue, readily incorporates into cells undergoing DNA synthesis. BrdU-pulsed cells are labeled with an enzyme-conjugated anti-BrdU antibody (Gratzner, H.G., 1982, Science 218: 474-475.). A colorimetric, soluble substrate is used to visualize proliferating cells that have incorporated BrdU. Reaction is stopped with sulfuric acid and plate is read at 450 nm using a microplate reader. Any statistically significant increase or decrease in color intensity of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function.

(F) Apoptosis by Annexin V

Programmed cell death or apoptosis is an early event in a cascade of catabolic reactions leading to cell death. A loss in the integrity of the cell membrane allows for the binding of fluorescently conjugated phosphatidylserine. Stained cells can be measured by fluorescence microscopy and flow cytometry (Vermes, I., 1995, J. Immunol. Methods. 180: 39-52). In one embodiment, any statistically significant increase or decrease in apoptotic cell number of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function.

For evaluating apoptosis in situ, assays for evaluating cell death in tissue samples can also be used in vivo studies.

(G) Quantitation of cytokine production

Cell supernatants harvested after cell stimulation for 16-48 hrs are stored at -80°C until assayed or directly tested for cytokine production. Multiple cytokine assays can be performed on each sample. IL-2, IL-3, IFN-γ and other cytokine ELISA Assays are available for mouse, rat, and human (Endogen, Inc. and BioSource). Cytokine production is measured using a standard two-antibody sandwich ELISA protocol as described by the manufacturer. The presence of horseradish peroxidase is detected with 3, 3', 5' tertamethyl benziidine (TMB) substrate and the reaction is stopped with sulfuric acid. The absorbency at 450 nm is measured using a microplate reader. Any statistically significant increase or decrease in color intensity of the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function. See also Example 1, *infra*. Detection of intracellular cytokines using anti-cytokine antibodies provides the additional advantage of measuring cytokines fore mixed cell populations. This allows for phenotyping measuring frequency of cytokine producing cell types in suspension or in tissues.

(H) NF-AT can be visualized by Immunostaining

T cell activation requires the import of nuclear factor of activated T cells (NF-AT) to the nucleus. This translocation of NF-AT can be visualized by immunostaining with anti-NF-AT antibody (Cell 1998, 93: 851-861). Therefore, NF-AT nuclear translocation has been used to assay T cell activation. Similarly, NF-AT/luciferase reporter assays have been used as a standard measurement of T cell activation (MCB 1996, 12: 7151-7160). Any statistically significant increase or decrease in the nuclear translocation of NF-AT brought about by the PDZ/PL interaction antagonist-treated sample, as compared to control sample (no treatment), suggest an effect of the PDZ/PL interaction antagonist on biological function. In order to optimize the use of the peptides and peptide analogues disclosed herein in a human subject, various animal models may be used to define certain clinical parameters. For example, the compounds of the invention may be tested in different dosages, formulations and route of administration in a cardiac transplant mouse model to optimize their ability to inhibit rejection responses to solid organ transplants (Fulmer et al., 1963, *Am. J. Anat.* 113:273; Jockusch et al., 1983, *Exp. Neurol.* 81:749).

In situations where inhibition of a T cell response is desired, the compounds of the inventions may be used to inhibit PDZ domain interactions with CD3, CD4, CD6 and CDw137. In addition, the compounds of the invention may be used to inhibit PDZ domain interactions with CD53 and CD138 in B cells. In order to inhibit IgE-mediated allergic reactions, the compounds of the invention may be used to inhibit PDZ domain interactions with FcεRIβ, CDw125 and CDw128. Furthermore, a PDZ motif sequence (PL sequence) of CD95 may be used to induce apoptosis of lymphomas.

(I) Inflammatory Mediator Release Assays

Assays are well known in the art for inflammatory mediator release to access the effect of compounds or treatments IgE-mediated degranulation. See, e.g. Berger et al., 1997, Measuring Cell Degranulation e.g., Ch 19.6 Immunology Method Manual. Academic Press, Ltd. 1436-1440 and Siraganian, 1983, Histamine Secretion from Mast Cells and Basophil. TIPS 4:432-437, both incorporated by reference herein.

6.9. Formulation and Route of Administration

6.9.1 Introduction of Agonists or Antagonists (e.g., Peptides and Fusion Proteins) into Cells

In one aspect, the PDZ-PL antagonists of the invention are introduced into a cell to modulate (i.e., increase or decrease) a biological function or activity of the cell. Many small organic molecules readily cross the cell membranes (or can be modified by one of skill using routine methods to increase the ability of compounds to enter cells, e.g., by reducing or eliminating charge, increasing lipophilicity, conjugating the molecule to a moiety targeting a cell surface receptor such that after interacting with the receptor). Methods for introducing larger molecules, e.g., peptides and fusion proteins are also well known, including, e.g., injection, liposome-mediated fusion, application of a hydrogel, conjugation to a targeting moiety conjugate endocytosed by the cell, electroporation, and the like).

In one embodiment, the antagonist or agent is a fusion polypeptide or derivatized polypeptide. A fusion or derivatized protein may include a targeting moiety that increases the ability of the polypeptide to traverse a cell membrane or causes the polypeptide to be delivered to a specified cell type (e.g., liver cells or tumor cells) preferentially or cell

compartment (e.g., nuclear compartment) preferentially. Examples of targeting moieties include lipid tails, amino acid sequences such as antennapodia peptide or a nuclear localization signal (NLS; e.g., *Xenopus* nucleoplasmin Robbins et al., 1991, *Cell* 64:615).

In one embodiment of the invention, a peptide sequence or peptide analog determined to inhibit a PDZ domain-PL protein binding, in an assay of the invention is introduced into a cell by linking the sequence to an amino acid sequence that facilitates its transport through the plasma membrane (a "transmembrane transporter sequence"). The peptides of the invention may be used directly or fused to a transmembrane transporter sequence to facilitate their entry into cells. In the case of such a fusion peptide, each peptide may be fused with a heterologous peptide at its amino terminus directly or by using a flexible polylinker such as the pentamer G-G-G-G-S (SEQ ID NO:1) repeated 1 to 3 times. Such linker has been used in constructing single chain antibodies (scFv) by being inserted between V_H and V_L (Bird et al., 1988, *Science* 242:423-426; Huston et al., 1988, *Proc. Natl. Acad. Sci. U.S.A.* 85:5979-5883). The linker is designed to enable the correct interaction between two beta-sheets forming the variable region of the single chain antibody. Other linkers which may be used include Glu-Gly-Lys-Ser-Ser-Gly-Ser-Gly-Ser-Glu-Ser-Lys-Val-Asp (SEQ ID NO:2) (Chaudhary et al., 1990, *Proc. Natl. Acad. Sci. U.S.A.* 87:1066-1070) and Lys-Glu-Ser-Gly-Ser-Val-Ser-Ser-Glu-Gln-Leu-Ala-Gln-Phe-Arg-Ser-Leu-Asp (SEQ ID NO:3) (Bird et al., 1988, *Science* 242:423-426).

A number of peptide sequences have been described in the art as capable of facilitating the entry of a peptide linked to these sequences into a cell through the plasma membrane (Derossi et al., 1998, *Trends in Cell Biol.* 8:84). For the purpose of this invention, such peptides are collectively referred to as transmembrane transporter peptides. Examples of these peptide include, but are not limited to, tat derived from HIV (Vives et al., 1997, *J. Biol. Chem.* 272:16010; Nagahara et al., 1998, *Nat. Med.* 4:1449), antennapedia from *Drosophila* (Derossi et al., 1994, *J. Biol. Chem.* 261:10444), VP22 from herpes simplex virus (Elliot and D'Hare, 1997, *Cell* 88:223-233), complementarity-determining regions (CDR) 2 and 3 of anti-DNA antibodies (Avrameas et al., 1998, *Proc. Natl. Acad. Sci. U.S.A.*, 95:5601-5606), 70 KDa heat shock protein (Fujihara, 1999, *EMBO J.* 18:411-419) and transportan (Pooga et al., 1998, *FASEB J.* 12:67-77). In a preferred embodiment of the invention, a truncated HIV tat peptide having the sequence of GYGRKKRRQRRRG (SEQ ID NO:173) is used.

It is preferred that a transmembrane transporter sequence is fused to a hematopoietic cell surface receptor carboxyl terminal sequence at its amino-terminus with or without a linker. Generally, the C-terminus of a PDZ motif sequence (PL sequence) must be free in order to interact with a PDZ domain. The transmembrane transporter sequence may be used in whole or in part as long as it is capable of facilitating entry of the peptide into a cell.

In an alternate embodiment of the invention, a hematopoietic cell surface receptor C-terminal sequence may be used alone when it is delivered in a manner that allows its entry into cells in the absence of a transmembrane transporter sequence. For example, the peptide may be delivered in a liposome formulation or using a gene therapy approach by delivering a coding sequence for the PDZ motif alone or as a fusion molecule into a target cell.

The compounds of the of the invention may also be administered via liposomes, which serve to target the conjugates to a particular tissue, such as lymphoid tissue, or targeted selectively to infected cells, as well as increase the half-life of the peptide composition. Liposomes include emulsions, foams, micelles, insoluble monolayers, liquid crystals, phospholipid dispersions, lamellar layers and the like. In these preparations the peptide to be delivered is incorporated as part of a liposome, alone or in conjunction with a molecule which binds to, e.g., a receptor prevalent among lymphoid cells, such as monoclonal antibodies which bind to the CD45 antigen, or with other therapeutic or immunogenic compositions. Thus, liposomes filled with a desired peptide or conjugate of the invention can be directed to the site of lymphoid cells, where the liposomes then deliver the selected inhibitor compositions. Liposomes for use in the invention are formed from standard vesicle-forming lipids, which generally include neutral and negatively charged phospholipids and a sterol, such as cholesterol. The selection of lipids is generally guided by consideration of, e.g., liposome size, acid lability and stability of the liposomes in the blood stream. A variety of methods are available for preparing liposomes, as described in, e.g., Szoka et al., Ann. Rev. Biophys. Bioeng. 9:467 (1980), U.S. Pat. Nos. 4,235,871, 4,501,728 and 4,837,028.

The targeting of liposomes using a variety of targeting agents is well known in the art (see, e.g., U.S. Patent Nos. 4,957,773 and 4,603,044). For targeting to the immune cells, a ligand to be incorporated into the liposome can include, e.g., antibodies or fragments thereof specific for cell surface determinants of the desired immune system cells. A liposome suspension containing a peptide or conjugate may be administered intravenously, locally, topically, etc. in a dose which varies according to, inter alia, the manner of administration, the

conjugate being delivered, and the stage of the disease being treated.

In order to specifically deliver a PDZ motif sequence (PL sequence) peptide into a specific cell type, the peptide may be linked to a cell-specific targeting moiety, which include but are not limited to, ligands for diverse leukocyte surface molecules such as growth factors, hormones and cytokines, as well as antibodies or antigen-binding fragments thereof. Since a large number of cell surface receptors have been identified in leukocytes, ligands or antibodies specific for these receptors may be used as cell-specific targeting moieties. For example, interleukin-2, B7-1 (CD80), B7-2 (CD86) and CD40 or peptide fragments thereof may be used to specifically target activated T cells (The Leucocyte Antigen Facts Book, 1997, Barclay et al. (eds.), Academic Press). CD28, CTLA-4 and CD40L or peptide fragments thereof may be used to specifically target B cells. Furthermore, Fc domains may be used to target certain Fc receptor-expressing cells such as monocytes.

Antibodies are the most versatile cell-specific targeting moieties because they can be generated against any cell surface antigen. Monoclonal antibodies have been generated against leukocyte lineage-specific markers such as certain CD antigens. Antibody variable region genes can be readily isolated from hybridoma cells by methods well known in the art.

However, since antibodies are assembled between two heavy chains and two light chains, it is preferred that a scFv be used as a cell-specific targeting moiety in the present invention. Such scFv are comprised of V_H and V_L domains linked into a single polypeptide chain by a flexible linker peptide.

The PDZ motif sequence (PL sequence) may be linked to a transmembrane transporter sequence and a cell-specific targeting moiety to produce a tri-fusion molecule. This molecule can bind to a leukocyte surface molecule, passes through the membrane and targets PDZ domains. Alternatively, a PDZ motif sequence (PL sequence) may be linked to a cell-specific targeting moiety that binds to a surface molecule that internalizes the fusion peptide.

In an other approach, microspheres of artificial polymers of mixed amino acids (proteinoids) have been used to deliver pharmaceuticals. For example, U.S. Pat. No. 4,925,673 describes drug-containing proteinoid microsphere carriers as well as methods for their preparation and use. These proteinoid microspheres are useful for the delivery of a number of active agents. Also see, U.S. Patent Nos. 5,907,030 and 6,033,884, which are incorporated herein by reference.

6.9.2 Introduction of Polynucleotides into Cells

A polynucleotide encoding a surface receptor C-terminal peptide may be useful in the treatment of various leukocyte activation-associated abnormal conditions. By introducing gene sequences into cells, gene therapy can be used to treat conditions in which leukocytes are activated to result in deleterious consequences. In one embodiment, a polynucleotide that encodes a PL sequence peptide of the invention is introduced into a cell where it is expressed. The expressed peptide then inhibits the interaction of PDZ proteins and PL proteins in the cell.

Thus, in one embodiment, the polypeptides of the invention are expressed in a cell by introducing a nucleic acid (e.g., a DNA expression vector or mRNA) encoding the desired protein or peptide into the cell. Expression may be either constitutive or inducible depending on the vector and choice of promoter. Methods for introduction and expression of nucleic acids into a cell are well known in the art and described herein.

In a specific embodiment, nucleic acids comprising a sequence encoding a peptide disclosed herein, are administered to a human subject. In this embodiment of the invention, the nucleic acid produces its encoded product that mediates a therapeutic effect by inhibiting leukocyte activation. Any of the methods for gene therapy available in the art can be used according to the present invention. Exemplary methods are described below.

For general reviews of the methods of gene therapy, see Goldspiel et al., 1993, Clinical Pharmacy 12:488-505; Wu and Wu, 1991, Biotherapy 3:87-95; Tolstoshev, 1993, Ann. Rev. Pharmacol. Toxicol. 32:573-596; Mulligan, 1993, Science 260:926-932; and Morgan and Anderson, 1993, Ann. Rev. Biochem. 62:191-217; May, 1993, TIBTECH 11(5):155-215. Methods commonly known in the art of recombinant DNA technology which can be used are described in Ausubel et al. (eds.), 1993, Current Protocols in Molecular Biology, John Wiley & Sons, NY; and Kriegler, 1990, Gene Transfer and Expression, A Laboratory Manual, Stockton Press, NY.

In a preferred embodiment of the invention, the therapeutic composition comprises a coding sequence that is part of an expression vector. In particular, such a nucleic acid has a promoter operably linked to the coding sequence, said promoter being inducible or constitutive, and, optionally, tissue-specific. In another specific embodiment, a nucleic acid molecule is used in which the coding sequence and any other desired sequences are flanked by regions that promote homologous recombination at a desired site in the genome, thus providing for intrachromosomal expression of the nucleic acid (Koller and Smithies, 1989, Proc. Natl. Acad. Sci. USA 86:8932-8935; Zijlstra et al., 1989, Nature 342:435-438).

Delivery of the nucleic acid into a patient may be either direct, in which case the patient is directly exposed to the nucleic acid or nucleic acid-carrying vector, or indirect, in which case, cells are first transformed with the nucleic acid *in vitro*, then transplanted into the patient. These two approaches are known, respectively, as *in vivo* or *ex vivo* gene therapy.

5 In a specific embodiment, the nucleic acid is directly administered *in vivo*, where it is expressed to produce the encoded product. This can be accomplished by any methods known in the art, *e.g.*, by constructing it as part of an appropriate nucleic acid expression vector and administering it so that it becomes intracellular, *e.g.*, by infection using a defective or attenuated retroviral or other viral vector (see U.S. Patent No. 4,980,286), by
10 direct injection of naked DNA, by use of microparticle bombardment (*e.g.*, a gene gun; Biolistic, Dupont), by coating with lipids or cell-surface receptors or transfecting agents, by encapsulation in liposomes, microparticles, or microcapsules, by administering it in linkage to a peptide which is known to enter the nucleus, or by administering it in linkage to a ligand subject to receptor-mediated endocytosis (see *e.g.*, Wu and Wu, 1987, J. Biol. Chem. 262:4429-
15 4432) which can be used to target cell types specifically expressing the receptors. In another embodiment, a nucleic acid-ligand complex can be formed in which the ligand comprises a fusogenic viral peptide to disrupt endosomes, allowing the nucleic acid to avoid lysosomal degradation. In yet another embodiment, the nucleic acid can be targeted *in vivo* for cell specific uptake and expression, by targeting a specific receptor (see, *e.g.*, PCT Publications WO
20 92/06180 dated April 16, 1992; WO 92/22635 dated December 23, 1992; WO92/20316 dated November 26, 1992; WO93/14188 dated July 22, 1993; WO 93/20221 dated October 14, 1993). Alternatively, the nucleic acid can be introduced intracellularly and incorporated within host cell DNA for expression, by homologous recombination (Koller and Smithies, 1989, Proc. Natl. Acad. Sci. USA 86:8932-8935; Zijlstra et al., 1989, Nature 342:435-438).

25 In a preferred embodiment of the invention, adenoviruses as viral vectors can be used in gene therapy. Adenoviruses have the advantage of being capable of infecting non-dividing cells (Kozarsky and Wilson, 1993, Current Opinion in Genetics and Development 3:499-503). Other instances of the use of adenoviruses in gene therapy can be found in Rosenfeld et al., 1991, Science 252:431-434; Rosenfeld et al., 1992, Cell 68:143-155; and
30 Mastrangeli et al., 1993, J. Clin. Invest. 91:225-234. Furthermore, adenoviral vectors with modified tropism may be used for cell specific targeting (WO98/40508). Adeno-associated virus (AAV) has also been proposed for use in gene therapy (Walsh et al., 1993, Proc. Soc. Exp. Biol. Med. 204:289-300).

In addition, retroviral vectors (see Miller et al., 1993, Meth. Enzymol. 217:581-
35 599) have been modified to delete retroviral sequences that are not necessary for packaging of

the viral genome and integration into host cell DNA. The coding sequence to be used in gene therapy is cloned into the vector, which facilitates delivery of the gene into a patient. More detail about retroviral vectors can be found in Boesen et al., 1994, *Biotherapy* 6:291-302, which describes the use of a retroviral vector to deliver the *mdr1* gene to hematopoietic stem cells in order to make the stem cells more resistant to chemotherapy. Other references illustrating the use of retroviral vectors in gene therapy are: Clowes et al., 1994, *J. Clin. Invest.* 93:644-651; Kiem et al., 1994, *Blood* 83:1467-1473; Salmons and Gunzberg, 1993, *Human Gene Therapy* 4:129-141; and Grossman and Wilson, 1993, *Curr. Opin. in Genetics and Devel.* 3:110-114.

Another approach to gene therapy involves transferring a gene to cells in tissue culture. Usually, the method of transfer includes the transfer of a selectable marker to the cells. The cells are then placed under selection to isolate those cells that have taken up and are expressing the transferred gene. Those cells are then delivered to a patient.

In this embodiment, the nucleic acid is introduced into a cell prior to administration *in vivo* of the resulting recombinant cell. Such introduction can be carried out by any method known in the art, including but not limited to transfection, electroporation, lipofection, microinjection, infection with a viral or bacteriophage vector containing the nucleic acid sequences, cell fusion, chromosome-mediated gene transfer, microcell-mediated gene transfer, spheroplast fusion, etc. Numerous techniques are known in the art for the introduction of foreign genes into cells (see *e.g.*, Loeffler and Behr, 1993, *Meth. Enzymol.* 217:599-618; Cohen et al., 1993, *Meth. Enzymol.* 217:618-644; Cline, 1985, *Pharmac. Ther.* 29:69-92) and may be used in accordance with the present invention, provided that the necessary developmental and physiological functions of the recipient cells are not disrupted. The technique should provide for the stable transfer of the nucleic acid to the cell, so that the nucleic acid is expressible by the cell and preferably heritable and expressible by its cell progeny. In a preferred embodiment, the cell used for gene therapy is autologous to the patient.

In a specific embodiment, the nucleic acid to be introduced for purposes of gene therapy comprises an inducible promoter operably linked to the coding sequence, such that expression of the nucleic acid is controllable by controlling the presence or absence of the appropriate inducer of transcription.

Oligonucleotides such as anti-sense RNA and DNA molecules, and ribozymes that function to inhibit the translation of a leukocyte surface receptor mRNA, especially its C-terminus are also within the scope of the invention. Anti-sense RNA and DNA molecules act to directly block the translation of mRNA by binding to targeted mRNA and preventing protein translation. In regard to antisense DNA, oligodeoxyribonucleotides derived from the

translation initiation site, *e.g.*, between -10 and +10 regions of a nucleotide sequence, are preferred.

The antisense oligonucleotide may comprise at least one modified base moiety which is selected from the group including, but not limited to, 5-fluorouracil, 5-bromouracil, 5-chlorouracil, 5-iodouracil, hypoxanthine, xanthine, 4-acetylcytosine, 5-(carboxyhydroxymethyl) uracil, 5-carboxymethylaminomethyl-2-thiouridine, 5-carboxymethylaminomethyluracil, dihydrouracil, beta-D-galactosylqueosine, inosine, N6-isopentenyladenine, 1-methylguanine, 1-methylinosine, 2,2-dimethylguanine, 2-methyladenine, 2-methylguanine, 3-methylcytosine, 5-methylcytosine, N6-adenine, 7-methylguanine, 5-methylaminomethyluracil, 5-methoxyaminomethyl-2-thiouracil, beta-D-mannosylqueosine, 5'-methoxycarboxymethyluracil, 5-methoxyuracil, 2-methylthio-N6-isopentenyladenine, uracil-5-oxyacetic acid (v), wybutoxosine, pseudouracil, queosine, 2-thiocytosine, 5-methyl-2-thiouracil, 2-thiouracil, 4-thiouracil, 5-methyluracil, uracil-5-oxyacetic acid methylester, uracil-5-oxyacetic acid (v), 5-methyl-2-thiouracil, 3-(3-amino-3-N-2-carboxypropyl) uracil, (acp3)w, and 2,6-diaminopurine.

Ribozymes are enzymatic RNA molecules capable of catalyzing the specific cleavage of RNA. The mechanism of ribozyme action involves sequence specific hybridization of the ribozyme molecule to complementary target RNA, followed by endonucleolytic cleavage. Within the scope of the invention are engineered hammerhead motif ribozyme molecules that specifically and efficiently catalyze endonucleolytic cleavage of leukocyte surface receptor RNA sequences.

Specific ribozyme cleavage sites within any potential RNA target are initially identified by scanning the target molecule for ribozyme cleavage sites which include the following sequences, GUA, GUU and GUC. Once identified, short RNA sequences of between 15 and 20 ribonucleotides corresponding to the region of the target gene containing the cleavage site may be evaluated for predicted structural features such as secondary structure that may render the oligonucleotide sequence unsuitable. The suitability of candidate targets may also be evaluated by testing their accessibility to hybridization with complementary oligonucleotides, using ribonuclease protection assays.

The anti-sense RNA and DNA molecules and ribozymes of the invention may be prepared by any method known in the art for the synthesis of nucleic acid molecules. These include techniques for chemically synthesizing oligodeoxyribonucleotides well known in the art such as for example solid phase phosphoramidite chemical synthesis. Alternatively, RNA molecules may be generated by *in vitro* and *in vivo* transcription of DNA sequences encoding the RNA molecule. Such DNA sequences may be incorporated into a wide variety of vectors

which contain suitable RNA polymerase promoters such as the T7 or SP6 polymerase promoters. Alternatively, antisense cDNA constructs that synthesize antisense RNA constitutively or inducibly, depending on the promoter used, can be introduced stably into cell lines.

5 Various modifications to the DNA molecules may be introduced as a means of increasing intracellular stability and half-life. Possible modifications include, but are not limited to, the addition of flanking sequences of ribo- or deoxy- nucleotides to the 5' and/or 3' ends of the molecule or the use of phosphorothioate or 2' O-methyl rather than phosphodiesterase linkages within the oligodeoxyribonucleotide backbone.

10 6.9.3 Other Pharmaceutical Compositions

 The compounds of the invention, may be administered to a subject *per se* or in the form of a sterile composition or a pharmaceutical composition. Pharmaceutical compositions comprising the compounds of the invention may be manufactured by means of
15 conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or lyophilizing processes. Pharmaceutical compositions may be formulated in conventional manner using one or more physiologically acceptable carriers, diluents, excipients or auxiliaries that facilitate processing of the active peptides or peptide analogues into preparations which can be used pharmaceutically. Proper formulation is
20 dependent upon the route of administration chosen.

 For topical administration the compounds of the invention may be formulated as solutions, gels, ointments, creams, suspensions, etc. as are well-known in the art.

 Systemic formulations include those designed for administration by injection, *e.g.* subcutaneous, intravenous, intramuscular, intrathecal or intraperitoneal injection, as well
25 as those designed for transdermal, transmucosal, oral or pulmonary administration.

 For injection, the compounds of the invention may be formulated in aqueous solutions, preferably in physiologically compatible buffers such as Hanks's solution, Ringer's solution, or physiological saline buffer. The solution may contain formulatory agents such as suspending, stabilizing and/or dispersing agents.

30 Alternatively, the compounds may be in powder form for constitution with a suitable vehicle, *e.g.*, sterile pyrogen-free water, before use.

 For transmucosal administration, penetrants appropriate to the barrier to be permeated are used in the formulation. Such penetrants are generally known in the art. This route of administration may be used to deliver the compounds to the nasal cavity.



For oral administration, the compounds can be readily formulated by combining the active peptides or peptide analogues with pharmaceutically acceptable carriers well known in the art. Such carriers enable the compounds of the invention to be formulated as tablets, pills, dragees, capsules, liquids, gels, syrups, slurries, suspensions and the like, for oral ingestion by a patient to be treated. For oral solid formulations such as, for example, powders, capsules and tablets, suitable excipients include fillers such as sugars, such as lactose, sucrose, mannitol and sorbitol; cellulose preparations such as maize starch, wheat starch, rice starch, potato starch, gelatin, gum tragacanth, methyl cellulose, hydroxypropylmethyl-cellulose, sodium carboxymethylcellulose, and/or polyvinylpyrrolidone (PVP); granulating agents; and binding agents. If desired, disintegrating agents may be added, such as the cross-linked polyvinylpyrrolidone, agar, or alginic acid or a salt thereof such as sodium alginate.

If desired, solid dosage forms may be sugar-coated or enteric-coated using standard techniques.

For oral liquid preparations such as, for example, suspensions, elixirs and solutions, suitable carriers, excipients or diluents include water, glycols, oils, alcohols, etc. Additionally, flavoring agents, preservatives, coloring agents and the like may be added.

For buccal administration, the compounds may take the form of tablets, lozenges, etc. formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently delivered in the form of an aerosol spray from pressurized packs or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurized aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of e.g. gelatin for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

The compounds may also be formulated in rectal or vaginal compositions such as suppositories or retention enemas, e.g., containing conventional suppository bases such as cocoa butter or other glycerides.

In addition to the formulations described previously, the compounds may also be formulated as a depot preparation. Such long acting formulations may be administered by implantation (for example subcutaneously or intramuscularly) or by intramuscular injection.

Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

Alternatively, other pharmaceutical delivery systems may be employed. Liposomes and emulsions are well known examples of delivery vehicles that may be used to deliver peptides and peptide analogues of the invention. Certain organic solvents such as dimethylsulfoxide also may be employed, although usually at the cost of greater toxicity.

5 Additionally, the compounds may be delivered using a sustained-release system, such as semipermeable matrices of solid polymers containing the therapeutic agent. Various of sustained-release materials have been established and are well known by those skilled in the art. Sustained-release capsules may, depending on their chemical nature, release the compounds for a few weeks up to over 100 days. Depending on the chemical nature and the
10 biological stability of the therapeutic reagent, additional strategies for protein stabilization may be employed.

As the compounds of the invention may contain charged side chains or termini, they may be included in any of the above-described formulations as the free acids or bases or as pharmaceutically acceptable salts. Pharmaceutically acceptable salts are those salts which
15 substantially retain the biologic activity of the free bases and which are prepared by reaction with inorganic acids. Pharmaceutical salts tend to be more soluble in aqueous and other protic solvents than are the corresponding free base forms.

6.10. Effective Dosages

The compounds of the invention will generally be used in an amount effective
20 to achieve the intended purpose. For use to inhibit leukocyte activation-associated disorders, the compounds of the invention or pharmaceutical compositions thereof, are administered or applied in a therapeutically effective amount. By therapeutically effective amount is meant an amount effective ameliorate or prevent the symptoms, or prolong the survival of, the patient being treated. Determination of a therapeutically effective amount is well within the
25 capabilities of those skilled in the art, especially in light of the detailed disclosure provided herein. An "inhibitory amount" or "inhibitory concentration" of a PL-PDZ binding inhibitor is an amount that reduces binding by at least about 40%, preferably at least about 50%, often at least about 70%, and even as much as at least about 90%. Binding can as measured *in vitro* (e.g., in an A assay or G assay) or *in situ*.

30 For systemic administration, a therapeutically effective dose can be estimated initially from *in vitro* assays. For example, a dose can be formulated in animal models to achieve a circulating concentration range that includes the IC₅₀ as determined in cell culture (*i.e.*, the concentration of test compound that inhibits 50% of leukocyte surface receptor-PDZ

domain-containing protein interactions). Such information can be used to more accurately determine useful doses in humans.

Initial dosages can also be estimated from *in vivo* data, *e.g.*, animal models, using techniques that are well known in the art. One having ordinary skill in the art could readily optimize administration to humans based on animal data.

Dosage amount and interval may be adjusted individually to provide plasma levels of the compounds that are sufficient to maintain therapeutic effect. Usual patient dosages for administration by injection range from about 0.1 to 5 mg/kg/day, preferably from about 0.5 to 1 mg/kg/day. Therapeutically effective serum levels may be achieved by administering multiple doses each day.

In cases of local administration or selective uptake, the effective local concentration of the compounds may not be related to plasma concentration. One having skill in the art will be able to optimize therapeutically effective local dosages without undue experimentation.

The amount of compound administered will, of course, be dependent on the subject being treated, on the subject's weight, the severity of the affliction, the manner of administration and the judgment of the prescribing physician.

The therapy may be repeated intermittently while symptoms detectable or even when they are not detectable. The therapy may be provided alone or in combination with other drugs. In the case of conditions associated with leukocyte activation such as transplantation rejection and autoimmunity, the drugs that may be used in combination with the compounds of the invention include, but are not limited to, steroid and non-steroid anti-inflammatory agents.

6.10.1 Toxicity

Preferably, a therapeutically effective dose of the compounds described herein will provide therapeutic benefit without causing substantial toxicity.

Toxicity of the compounds described herein can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, *e.g.*, by determining the LD₅₀ (the dose lethal to 50% of the population) or the LD₁₀₀ (the dose lethal to 100% of the population). The dose ratio between toxic and therapeutic effect is the therapeutic index. Compounds which exhibit high therapeutic indices are preferred. The data obtained from these cell culture assays and animal studies can be used in formulating a dosage range that is not toxic for use in human. The dosage of the compounds described herein lies preferably within a range of circulating concentrations that include the effective dose with little or no toxicity.

The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. The exact formulation, route of administration and dosage can be chosen by the individual physician in view of the patient's condition. (See, e.g., Fingl *et al.*, 1975, In: The Pharmacological Basis of Therapeutics, Ch.1, p.1).

7. EXAMPLES

7.1 EXAMPLE 1: TAT-T CELL SURFACE RECEPTOR CARBOXYL TERMINUS FUSION PEPTIDES INHIBITED T CELL ACTIVATION

7.1.1. Materials And Methods

7.1.1.1. Peptide Synthesis

All peptides were chemically synthesized by standard procedures. The Tat-CD3 carboxyl terminus fusion peptide, (GYGRKKRRQRRRGPPSSSSGL, SEQ ID NO:174); Tat-CLASP1 carboxyl terminus fusion peptide, (GYGRKKRRQRRRGSISSSAEV, SEQ ID NO:243); Tat-CLASP2 carboxyl terminus fusion peptide, (GYGRKKRRQRRRGMTSSSSVV, SEQ ID NO:176); and Tat peptide, (GYGRKKRRQRRRG, SEQ ID NO:173); were dissolved at 1 mM in PBS, pH 7, or dH₂O. Stock MBPac1-16 peptide, (AcASQKRPSQRHGSKYLA, SEQ ID NO: 403), was dissolved at 5 mM. All peptides were aliquoted and stored at -80°C until tested.

7.1.1.2 Cell Cultures

Cells were maintained and tested in RPMI 1640 media supplemented with 10% fetal calf serum (HyClone), 2 mM glutamine, 10 mM Hepes, 100 U/ml penicillin, 100 µg/ml streptomycin, 0.1 mM non-essential amino acids, 1 mM sodium pyruvate, and 50 µM beta mercaptoethanol.

7.1.1.3 T Cell Stimulation Assay

Supernatants were assayed for cytokine production following activation of T cell lines. Mouse T cell lines were stimulated using two different methods, either with antigen and antigen presenting cells or anti-mouse CD3.

Antigen-specific mouse T cells, BR4.2, were activated with the N-terminal 16 amino acid sequences of myelin basic protein (MBPac1-16) and syngenic mouse splenocytes in 96-well plates. Mitomycin C-treated antigen presenting cells, 2×10^5 B10.BR, were added to each row of serially diluted MBPac1-16 ranging from 0 to 200 µM. Next, 10 µM Tat-peptides or media alone was added to each row. Finally, 2×10^4 MBPac1-16-specific T cell, pre-loaded with 10 µM Tat-peptides (see above), were added to all wells (Rabinowitz et al., 1997, Proc. Natl. Acad. Sci. U.S.A., 94:8702-8707). Cells were activated during an

overnight incubation at 5% CO₂, 37⁰C. Cell supernatant was collected and stored at -80⁰C until assayed for cytokine production. The final volume was 200 µl/well.

Antibody against mouse CD3 (Pharmigen #145-2C11) was coated overnight at 4⁰C using 96-well flat bottom Elisa plates at a final concentration of 0.5 µg/ml, diluted in PBS.

5 Just prior to use, plates were washed three times with 200 µl/well PBS to remove excess anti-CD3. To ensure that cells were given sufficient time to transduce Tat-peptides before activation, T cells (5x10⁵ cells/ml) were pre-treated with or without 10 µM Tat-peptides for two hours at 5% CO₂, 37⁰C and then diluted in media with or without 10 µM Tat-peptides to a final concentration of 2x10⁴ cells per well in a final volume of 200 µl. Cells were then treated as
10 described above.

7.1.1.4 Cytokine ELISA

IFN γ was measured from cell supernatants, described above, at ambient temperature using the Endogen, Inc. ELISA protocol 3. Briefly, 96-well, flat bottom, high
15 binding ELISA plates were preincubated overnight with coating antibody (MM700). Plates were washed with 50 mM TRIS, 0.2% tween-20, pH 8 and they blocked for one hour with PBS plus 2% BSA. Washed plates were then incubated one hour with 25 µl of cell supernatant and 25 µl blocking buffer, or with 50 µl IFN γ standard. The presence of IFN γ was detected with a biotin-labeled anti-mouse IFN γ monoclonal antibody (MM700B, Endogen, Inc.,).

20 Quantitative amounts of detection antibody are revealed with horseradishperoxidase-conjugated streptavidin. The enzymatic, color, substrate for HRP, tetramethylbenzidine (TMB), was developed for up to 30 minutes and stopped with 1.0 M H₂SO₄. The absorbance at 450 nm was measured using a microtiter plate reader (Thermo Max, Molecular Devices) and the concentration of unknown IFN γ from cell supernatants was calculated from a standard curve
25 generated by Softmax Pro. software (Molecular Devices).

7.1.1.2 Results

Peptides containing Tat transporter sequences linked to C-terminal sequences of various PLs were testing for their ability to inhibit T cell activation. **FIGURE 4A** shows
30 that the Tat-CD3 fusion peptide inhibits T cell activation mediated by peptide:MHC as compared to controls of Tat-peptide alone or no peptide. **FIGURE 4B** shows that Tat-CLASP2 carboxyl terminus fusion peptide inhibited T cell activation mediated by monoclonal anti-CD3

as compared to Tat-peptide alone. Tat-CLASP1 fusion peptide did not inhibit T cell activation in this experiment. These results indicate that peptides containing potential inhibitory sequences can be transported into T cells through transporter peptide such as Tat to disrupt surface receptor organization mediated by PDZ proteins. Disruption of PDZ-mediated surface
5 receptor organization leads to blockage of T cell activation in response to antigen.

7.2. EXAMPLE 2: DESIGN OF AN INHIBITOR OF DLG 1-LIGAND BINDING WITH GREATER THAN 100 uM POTENCY

10 A GST/DLG1 fusion protein (See TABLE 3) and a biotin-labeled peptide corresponding to the C-terminal 20 amino acids of the CLASP-2 protein, peptide AA2L (see TABLE 4), were synthesized and purified by standard techniques well known in the art as described supra. This PDZ-ligand combination was then shown to bind specifically using both the “A” assay and the “G” assay (See TABLE 2). Once specific binding was demonstrated, the
15 apparent affinity of the binding interaction was determined using Approach 1 of the section entitled “Measurement of PDZ-ligand binding affinity” (see FIGURE 2A). The measured apparent affinity was 21 uM. This implies that 21 uM labeled CLASP-2 peptide AA2L filled 50% of the binding sites for CLASP-2 on DLG1. Thus, 21 uM unlabeled CLASP-2 peptide should be able to block the binding of a given ligand to DLG1 by approximately 50%, assuming
20 that the given ligand (1) binds to the same site(s) on DLG1 as Qasp~2 and (2) is not added at sufficient concentration to reduce significantly the binding of the CLASP-2 peptide (i.e. cannot out-compete the CLASP-2 peptide).

To detect such inhibition, it was necessary to synthesize an analogue of the CLASP2peptide AA2L that (1) retained similar DLG1 binding properties and (2) would not
25 itself generate a signal in the assay selected to measure inhibition. Because most molecular interactions between PDZ proteins and their ligands involve only the C-terminal 6 amino acids of the ligand, an eight amino acid variant of the CLASP-2 peptide, MTSSSSVV (SEQ ID NO: 191), was anticipated to retain similar DLG1 binding properties as the 20 amino acid AA2L CLASP-2 peptide. This eight amino acid CLASP-2 peptide (lacking a functional label) was
30 therefore synthesized and purified by standard techniques as described *supra*. When 100 uM of the (functionally unlabeled) eight amino acid CLASP-2 peptide and 20 uM of the biotin-labeled AA2L CLASP-2 peptide were added simultaneously to DLG1 in a variant of the “G” assay (described supra), the binding of the labeled AA2L CLASP-2 peptide was, as predicted,

inhibited by greater than 50% (**FIGURE 3A**). An analogous experiment in which the labeled AA2L CLASP-2 peptide was replaced with another labeled DLG1 ligand, labeled AAI3L Fas peptide demonstrated similar inhibition by the eight amino acid CLASP-2 peptide (**FIGURE 3A**). Thus, an effective inhibitor of DLG1-ligand binding (i.e. the eight amino acid CLASP-2 peptide MTSSSSVV, SEQ ID NO: 191) with a known potency range (order of magnitude 21 uM) was designed based on knowledge of the affinity, 21 uM, with which a particular labeled ligand, the CLASP-2 peptide AA2L, bound to DLG1.

7.3 **EXAMPLE 3:** GENERATION OF EUKARYOTIC EXPRESSION CONSTRUCTS BEARING DNA FRAGMENTS THAT ENCODE PDZ DOMAIN CONTAINING GENES OR PORTIONS OF PDZ DOMAIN GENES

This example describes the cloning of PDZ domain containing genes or portions of PDZ domain containing genes into eukaryotic expression vectors in fusion with red fluorescent protein (RFP).

A. Strategy

DNA fragments corresponding to PDZ domain containing genes were generated by RT-PCR from jurkat cell line (transformed T-cells) derived RNA. Primers were designed to create restriction nuclease recognition sites at the PCR fragment's ends, to allow cloning of those fragments into the appropriate vectors. Subsequent to RT-PCR, DNA samples submitted to agarose gel electrophoresis. Bands corresponding in size to the expected size were excised, DNA extracted and treated with appropriate restriction endonuclease. DNA samples were purified once more by gel electrophoresis, and gel extracted DNA fragments were coprecipitated and ligated with the appropriate linearized cloning vector. After transformation into *E.coli*, bacterial colonies were screened by PCR for the presence and correct orientation of insert. Positive clones were picked for large scale DNA preparation and the insert including the flanking vectors sites were sequenced to ensure correct sequence of fragments and junctions with the vectors and fusionproteins.

B. Vectors:

Cloning vectors were pDsRED1-N1 (purchased from CLONTECH, # 6921-1) and pDsRED1-N1(+ATG), a derivative of pDsRED1-N1 generated by recombinant DNA technology.

DNA fragments to clone that contained the ATG-start codon were cloned into pDsRED1-N1. Fragments void of a proper translation initiation codon were cloned into pDsRED1-N-(+ATG), since this vector includes an translation initiation start codon. Vector pDsRED1-N1(+ATG) differs from pDsRED1 only with regard to the multiple cloning sites.

5 The sequence that is unique to pDsRED1-N1(+ATG) is shown below; boundaries with pDsRED1-N1 are printed in lower case and correspond to nucleotides N 633 and N 662 in pDsRED1-N1, respectively.

5'-attGCCACCATGGGAATTCTGGATCCGGGAgat-3'

10 C. Deduced amino acid linker sequences:

Linker sequences between the cloned inserts and RFP vary depending on the vectors and on the restriction endonuclease used for cloning. Deduced linker amino acid sequences are listed in the table below; For some constructs, the first N-terminal and / or last C-terminal amino acid corresponds to a linker amino acid introduced by the cloning process

15 but is not represented at that position in the corresponding gene.

Table 8

pDsRED1-N1, cloning approach: (fragment) Eco RI or Mfe I / Eco RI (vector)	PDZ domain insert C-term - LEU - GLN - SER - THR - VAL - PRO - ARG - ALA - ARG - ASP - PRO - PRO - VAL - ALA - THR - red flourecent protein;
pDsRED1-N1(+ATG), cloning approach: (fragment) Eco RI / Eco RI (vector)	Start codon (MET) - GLY - ILE - PDZ domain gene insert - LEU - ASP - PRO - GLY - TYR - PRO - PRO - VAL - ALA - THR - red flourecent protein;
pDsRED1-N1(+ATG), cloning approach: (fragment) Mfe I / Eco RI (vector)	Start codon (MET) - ARG - ILE - PDZ domain gene insert - LEU - ASP - PRO - GLY - TYR - PRO - PRO - VAL - ALA - THR - red flourecent protein;

20 D. Constructs:

The deduced protein sequence of cloned inserts, primers used to generate DNA fragments by RT-PCR and accession # are given below for each construct. For all constructs, the fusion with RFP was carboxy terminal.

25 1. Homo sapiens dishevelled 1 (DVL1)

Acc #:

NM_004421

GI: 4758213

Cloning sites for all constructs: Eco RI / Eco RI

- Construct (N-P) [Covers the methionin start codon and extends over the C-terminal

boundary of the DVL1 PDZ domain];

primers: 308 DVF and 311 DVR;

vector: pDsRED1-N1

5

aa 1 - aa 341

MAETKIIYHMDEEETPYLVKLPVAPERVTLADFKNVLSNRPVHAYKFFKSMDQDFGVV
KEEIFDDNAKLPCFNRRVSWLVLEGAHSDAGSQGTDSHTDLPPPLERTGGIGDSRSP
SFQPDVASSRDGMDNETGTESMVSHRRDRARRRNREEAARTNGHPRGDRRRDVGLPP
10 DSASTALSSELESSSFVDSDEDDSTSRLSSSTEQSTSSRLIRKHKRRRRKQRLRQADRAS
SFSSMTDSTMSLNITVTNLNMRHHFLGICIVGQSNDRGDGGIYIGSIMKGGAVAADGRI
EPGDMLLQVNDVNFENMSNDDAVRVLREIVSQTGPISLTVAKCWDPT

15

- Construct (N) [Covers the methionin start codon and extends to the N-terminal

boundary of the DVL1 PDZ domain];

primers: 308 DVF and 345 DVR

vector: pDsRED1-N1

20

aa 1 - aa 197

MAETKIIYHMDEEETPYLVKLPVAPERVTLADFKNVLSNRPVHAYKFFFKSMDQDFGV
VKEEIFDDNAKLPCFNRRVSWLVLEGAHSDAGSQGTDSHTDLPPPLERTGGIGDSRS
PSFQPDVASSRDGMDNETGTESMVSHRRDRARRRNREEAARTNGHPRGDRRRDVGLP
PDSASTALSSELESSSFVDSDEDG

25

- Construct (P) [Consists of the PDZ domain of DVL1];

primers: 344 DLF and 311 DVR;

vector: pDsRED1-N1(+ATG)

30

aa 246 - aa 341

SLNIITVTNLNMRHHFLGICIVGQSNDRGDGGIYIGSIMKGGAVAADGRIEPGDMLLQVN
DVNFENMSNDDAVRVLREIVSQTGPISLTVAKCWDPT

Primers:

35

308 DVF (N 128 - N 155) 5'-TCGGAATTCGTCGCGCCATGGCGGAGAC-3'

311 DVR (N 1004 - N 1032) 5'-GGGAATTCGGTCCCAGCACTTGGCCACAG-3'

344 DVF (N 873 - N 900) 5'-CCAGAATTCTCAACATCGTCACTGTAC-3'

345 DVR (N713 - N744) 5'-TCGGAATTCATCCTCGTCCGAGTCCACAAAG-3'

40

2. KIAA 0751 / 41.8 KD

Acc #:

AB018294

GI: 3882222

5

Cloning sites for all constructs: (vector) Eco RI / (fragment) Mfe I

• Construct (N-J) [includes the third in frame-methionin (putative start) codon in (GI: 3882222) and extends c-terminal of the PDZ domain to the region on sequence divergency between KIAA 0751 (GI: 3882222) and hypothetical 41.8 Kd protein (AF007156 / GI: 3882222)];

primers: 318 KIF and 320 KIR;

vector: pDsRED1-N1

aa 389 - aa 803

15 MMYFGGHSLEEDLEWSEPQIKDSGVDTCSSITLNEEHSKHPVTWQPSKDGDRDIG
RILLNKRLKDGSVPRDSGAMLGLKVVGKMTESGRLCAFITKVKKGSLADTVGHLRPG
DEVLEWNGRLLQGATFEEVYNILESKEPEQVELVVSRIPIGDIPIPDSTHAQLESSSSFE
SQKMDRPSISVTSPMSPGMLRDVPQFLSGQLSIKLWFDKVGHQLIVTILGAKDLPSRED
GRPRNPYVKIYFLPDRSDKNKRRTKTVKKTLEPKWNQTFIYSPVHRREFRERMLEITLW
20 DQARVREEESEFLGEILIELETALLDDEPHWYKLQTHDVSSLPLPHPSPYMPRRQLHGES
PTRRLQRSKRISDSEVSDYDCDDGIGVVSDYRHDGRDLQSSTLSVPEQVMSSNHCSPSG
SPHRVDVIGRTT

25 •Construct (P) [consists of the PDZ domain of KIAA 0751 / 41.8 Kd hypothetical protein (GI: 3882222)];
primers: 341 KIF and 319 KIR.
vector pDsRED1-N1(+ATG)

30 aa 443 - aa 534

LKDGSVPRDSGAMLGLKVVGKMTESGRLCAFITKVKKGSLADTVGHLRPGDEVLEW
NGRLLQGATFEEVYNILESKEPEQVELVVSRIPIA

Primers:

35 318 KIF (N 1366 - N 1393) 5'-AGACAATTGAGGAAATGATGTACTTTGG-3'
319 KIR (N 1830 - N 1857) 5'-GAACAATTGCAATAGGCCTTGAAACTAC-3'
320 KIR (N 2640 - N 2667) 5'-ACCCAATTGTAGTCCTTCCTATAACATC-3'
341 KIF (N 1567 - N 1593) 5'-ATAGAATTCTAAAAGATGGAAGTGTAC-3'

40 3. Homo sapiens PAR6

Acc #:

AF265565

GI: 8468608

5 Cloning sites for all constructs: Eco RI / Eco RI

- Construct (N-P) [Covers the methionin start codon and extends over the C-terminal boundary of the PDZ domain];

primers: 322 PAF and 324 PAR;

vector: pDsRED1-N1

10

aa 1 - aa 251

MARPQRTPARSPDSIVEVKSFKDAEFRRFALPRASVSGFQEFsRLLRAVHQIPGLDVLLG
YTDAHGDLPLTNDDSLHRALASGPPPLRLLVQKREADSSGLAFASNSLQRRKKGLLLR
PVAPLRTRPPLLISLPQDFRQVSSVIDVDLLPETHRRVRLHKHGS DRPLGFYIRDGMSVR
15 VAPQGLERVPGIFISRLVRGGLAESTGLLAVSDEILEVNGIEVAGKTLDQVTDMMVANS
HNLIVTVKPANQR

- Construct (N) [Covers the methionin start codon and extends to the N-terminal boundary of the PDZ domain];

20 primers: 322 PAF and 343 PAR

vector: pDsRED1-N1

aa 1 - aa 147

MARPQRTPARSPDSIVEVKSFKDAEFRRFALPRASVSGFQEFsRLLRAVHQIPGLDVLLG
25 YTDAHGDLPLTNDDSLHRALASGPPPLRLLVQKREADSSGLAFASNSLQRRKKGLLLR
PVAPLRTRPPLLISLPQDRQVSSVIDV

- Construct (P) [Consists of the PDZ domain of PAR6];

primers: 342 PAF and 324 PAR;

30 vector: pDsRED1-N1(+ATG)

aa 155 - aa 251

RRVRLHKHGS DRPLGFYIRDGMSVRVAPQGLERVPGIFISRLVRGGLAESTGLLAVSDEI
35 LEVNGIEVAGKTLDQVTDMMVANS HNLIVTVKPANQR

Primers

322 PAF (N 55 - N 82) 5'-CCCGAATTCGCCATGGCCCGGCCGCAGAG-3'

324 PAR (N 798 - N 825) 5'-CGTGAATTCGCTGGTTGGCGGGCTTGAC-3'

342 PAF (N 519 - N 548) 5'-GAGGAATTCCGACGGGTGCGGCTGCACAAG-3'

343 PAR (N 485 - N 516) 5'-GCAGAATTCCCACGTCTATGACTGAGGAAAC-3'

4. Homo sapiens post-synaptic density protein 95 (PSD95)

Acc #:

5 ABU83192

GI: 3318652

Cloning sites for all constructs: Eco RI / Eco RI

Vector: pDsRED1-N1

- Construct (N-P3) [Covers the methionin start codon and extends over the C-terminal

10 boundary of PDZ domain 3;

primers: 315 PSF and 304 PSR.

aa 1 - aa 442

15 MSQRPRAPRSALWLLAPPLLRWAPPLLTVLHSDLFQALLDILDYYEASLSESQKYRYQD
EDTPPLEHSPAHLPNQANSPPVIVNTDTLEAPGYELQVNGTEGEMEYEEITLERGNSGL
GFSIAGGTDNPHIGDDPSIFITKIIPGGAAAQDGRRLRVNDSILFVNEVDVREVTHSAAVEA
LKEAGSIVRLYVMRRKPPAEKVMEIKLIKGPGLGFSIAGGVGNQHIPGDNSIYVTKIIE
GGAHKDGRLLQIGDKILAVNSVGLDVMHEDAVAALKNTYDVVYLKVAKPSNAYLS
DSYAPPDITTSYSQHLNEISHSSYLGTDYPTAMTPTSPRRYSPVAKDLLGEEDIPREPR
20 RIVIHGSGTGLGFNIVGGEDGEGIFISFILAGGPADLSGELRKGDQILSVNGVDLRNASHE
QAAIALKNAGQTVTIIAQYKPEEYSR

primers:

315 PSF (N847 - N876) 5'-AGAGAATTCAGAGATATGTCCCAGAGACCAAG-3'

25 304 PSR (N 2161 - N 2189) 5'-CGAGAATTCTGTACTCTTCTGGTTTATAC-3'

5. Homo sapiens hCASK (CASK)

Acc #:

AF032119

30 GI: 2641548

Cloning sites: Eco RI / Eco RI

- Construct (P) [Covers the PDZ domain of hCASK];

Note: The amino acid sequence homology between the human hCASK and the mouse mCASK-B is 100% identical.

35 primers: 336 CAF and 335 CAR;

vector: pDsRED1-N1(+ATG)

aa 399 - aa 572

RLVQFQKNTDEPMGITLKMNELNHCIVARIMHGGMIHRQGLHVGDEIREINGISVANQ
TVEQLQKMLREMRGSITFKIVPSYR

5

Primers

336 CAF (N 1484 - N 1512) 5'-CCAGAATTCGGCTGGTACAGTTTCAAAAG-3'

325 CAR (N 1722 - N 1750) 5'-ACTGAATTCGGTAACTTGGCACAATCTTG-3'

10 6. Homo sapiens membrane protein, palmitolated 2 (MPP2 / DLG2)

Acc #:

X82895

GI: 939884

15 Cloning sites for all constructs: Eco RI / Eco RI

• Construct (N-SH3) [Covers the methionin start codon, the PDZ domain and extends to the C-terminal boundary of the MPP2 SH3 domain;the construct is a splice variant of the construct annotated under GI:939884. With respect to GI:939884, the DNA portion N 238 to 309 is missing; this DNA stretch corresponds to AA 51-74. The open reading frame is maintained throughout the deletion].

primers: 305 MF and 306 MR;

vector: pDsRED1-N1

aa 1 - aa 317

25 MPVAATNSETAMQQVLDNLGSLPSATGAAELDLIFLRGIMESPIVRSLAKAHERLEETK
LEAVRDNNLELVQEILRDLAQLAEQSSTAELAHLQEPHFQSLETHDSVASKTYETPP
PSPGLDPTFSNQPVPDAVRMVGIRKTAGEHLGVTFRVEGGELVIARILHGGMVAQQG
LLHVGDIKEVNGQPVGSDPRALQELLRNASGSVILKILPSYQEPHLPRQVFVKCHFDYD
PARDSLIPCKEAGLRFNAGDLLQIVNQDDANWWQACHVEGGSAGLIPSQLLEEKRKG

30

Primers:

305 MF (N 58 - N 84) 5'-AGAGAATTCAGAGCCCTTGCCTCCTTC-3'

306 MR (N 798 - N 825) 5'-TGAGAATTCCTTTCCGCTTCTCCTCCAG-3'

35 7. Homo sapiens Tax interaction protein 1 (TIP1)

Acc #:

AF028823

GI: 2613001

Cloning sites: Eco RI / Bam H1

(We determined 5' start site and 5' full length sequence by 5' RACE)

- Construct (N-C);

5 vector: pDsRed1-N1

aa 3 - aa 125

YIPGQPVTAVVQRVEIHKLRQGENLILGFSIGGGIDQDPSQNPFSEDKTDKGIYVTRVSE
GGPAEIAQLQSGDKIMQVNGWDMTMVTHDQARKRLTKRSEEVVRLLVTRQSLQKAV
10 QQSML

Primer:

1318 TIP R3-1 (N 336 - N 356) 5'-CAGTCCATGCTGTCGGATCCG-3'

1317 TIP R5-1* 5'-GTCGGAATCCCTACATCCCG-3'

15 *Primer 5' end corresponds to the nucleotide that is located 29 nucleotides 5' of N 1; primer sequence corresponds to sequence determined by 5' RACE; numbering corresponds to genbank sequence entry (GI 2613001).

7.4 EXAMPLE 4: EXPRESSION OF PDZ DOMAIN-CONTAINING PROTEINS IN MAMMALIAN CELLS

We have identified a number of PDZ domain-containing proteins that are expressed in lymphocytes. To study the biology of these molecules in immunity, we have adopted a dominant-negative approach by over-expressing portions of tagged PRISM molecules in lymphocytes. We have termed cellular studies of gene function as CELLOMICS. We used two standard methods for DNA transfection; DNA precipitation by calcium phosphate (Graham and van der Eb, *infra.* and Gorman,) and electroporation (Potter, *infra.*) Expression of PDZ fusion proteins was tested in mouse and human lymphocytic cell lines and in human embryonic kidney cells (293 HEK, ATCC CRL-1573) based on detection of red fluorescent protein (RFP).

7.4.1 Materials and Methods

A. Cell lines used for transfection of PRISM-tagged constructs

Jurkat E6 human T cells (ATCC TIB-152) were maintained and tested in complete IMDM (IMDM medium supplemented with 2 mM glutamine, 100 U/mL penicillin, 100 µg/mL streptomycin, 0.1 mM nonessential amino acids, 1 mM sodium pyruvate (Gibco BRL), 50 µM beta mercaptoethanol (Sigma), and 10% fetal calf serum (Gemini Bio-Products)). 293 HEK cells were maintained and tested in complete DMEM (DMEM medium supplemented with 2 mM glutamine, 10 mM HEPES, 100 U/mL penicillin, 100 µg/mL streptomycin, and 10% fetal calf serum). CH27 mouse B cell lymphoma and 2B4 mouse T cell hybrid lines were maintained and tested in complete RPMI (RPMI 1640 medium supplemented with 2 mM glutamine, 100 U/mL penicillin, 100 µg/mL streptomycin, 0.1 mM nonessential amino acids, 1 mM sodium pyruvate, 10 mM HEPES, 50 µM beta mercaptoethanol, and 10% fetal calf serum).

B. Transfection Methods

DNA precipitation by calcium phosphate was used to transfect 293 HEK cells. At least 2 hrs before transfection, $\sim 5 \times 10^5$ cells were plated in 5 ml of complete DMEM onto one 60 mm TC-treated dish for each transfection. For each plate, 5-10 µg of DNA was brought to a volume of 110 µl with deionized H₂O. Fifteen µl of 2M CaCl₂ was added to the DNA solution, which was then slowly added to 125 µl of 2X HBS pH 7.0, (2X HBS = 1.64% NaCl 1.188 % and Hepes 0.04% Na₂HPO₄). A fine precipitate formed, which was then added

dropwise to cells and incubated at 37°C, 5% CO₂. Transfected cells were analyzed for expression of RFP on day 1, 2 and 3 post-transfection using the RFP control plasmid and some of the PRISM-RFP fusion constructs. Since we consistently found maximal expression on day 2, the data described below is from cells 2 days after transfection.

5 The lymphocytic cell lines (Jurkat E6, CH27, and 2B4) cells were transfected by electroporation and tested for control RFP expression on day 1 and 2 post-transfection. All three lymphocytic cell lines showed that maximal expression of RFP control plasmid and some of the PRISM-RFP constructs on day 1 post-transfection and therefore only day 1 expression data is described below. The BTX ECM830 generator was used to transfect the Jurkat E6 cells as follows: two cuvettes (4 mm electrode gap) containing 5 x 10⁶ cells in 0.5 ml and 5-30 µg of DNA in serum free-IMDM were electroporated with a single pulse at 260 volts for 50 msec. Cuvettes were immediately placed on ice for 10-15 minutes before being transferred to 100 mm dishes containing 10 ml of complete IMDM. CH27 and 2B4 cells were transfected in cuvettes (4 mm electrode gap) using the BioRad Gene pulser. The protocol using the BioRad Gene pulser is the same as for the BTX ECM830 generator, except they were electroporated in serum free-RPMI at 0.45kV, 960 µF with unlimited resistance for 38-44 msec.

7.4.2 Analysis of PRISM-tagged fusion protein expression

A. PDZ-RFP fusion protein constructs

20 DNA fragments encoding PDZ domain proteins were cloned into DsRED (Clontech), an RFP fusion vector. To ensure proper folding of inserts, the RFP was placed C-terminal to PRISM. Constructs tested for expression in the DsRED vector are: CASK(P), DLG1(N-P3), DLG1(N), DVL1(N-P), DVL1(N), DVL1(P), MPP2(N-SH3), PAR6(N-P), PAR6(N), PAR6(P), PSD95(N-P3), TIP1(P), TIP1(N-C), KIAA0751(N3-J) and KIAA0751(P), as described in Example 3, *supra*. The abbreviation N denotes the N-terminus, C denotes the C-terminus, P denotes one PDZ region, P3 denotes three PDZ regions, SH3 denotes the inclusion of the SH3 domain, and J denotes the joining region.

B. PDZ-RFP fusion proteins are expressed in 293 HEK Cells

30 All of the PDZ-RFP transfected cells were analysis using a Coulter EPICS XL flow cytometer and inverted Nikon Diaphot fluorescent microscope. Each set of transfections included the DsRED plasmid (RFP) as a positive control for transfection efficiency. All of the

PDZ-RFP constructs expressed RFP fusion protein in 293 HEK cells with transfection efficiency ranging from 15-60%.

Fluorescent microscopy revealed various expression patterns. The RFP expression from the control DsRED construct was evenly diffused in the cell varying from weak to very brightly fluorescence. For the constructs expressed in 293 HEK cells, the staining patterns are summarized here: CASK(P) some bright and diffuse and some punctated, DLG1(N-P3) differentially speckled, some globular and few diffuse, DLG1(N) differentially speckled and some globular, DVL1(N-P) diffuse and punctated and polarized and globular, DVL1(N) brightly speckled and few polarized and globular, DVL1(P) occasional diffuse and rare speckled, MPP2(N-SH3) polarized and globular and some punctated, PAR6(N-P) speckled and some polarized and globular, PAR6(N) diffuse and punctated, PAR6(P) diffuse and punctated, polarized and globular, PSD95(N-P3) diffuse, TIP1(P) diffuse, speckled, polarized and globular, TIP1(N-C) occasional rings and polarized, KIAA0751(N3-J) punctated and few dim and diffuse, and KIAA0751(P) punctated and few dim and diffuse.

C. PDZ-RFP fusion protein expression is restricted in Jurkat T cells

Jurkat E6 cells were screened for expression of PDZ-RFP fusion proteins tested using flow cytometry and a fluorescent microscope on day 1 post-transfection. Although expression of all in 293 HEK cells, several constructs did not expressed in Jurkat E6 cells. The control RFP and all of the expressed PDZ-RFP fusion proteins had a much lower percent of cells expressing RFP as compared to 293 HEK cells. Constructs that were tested but less than 4% of the cells expressed RFP were considered negative, they were: DLG1(N-P3), DLG1(N), DVL1(P), MPP2(N-SH3), PAR6(N-C), PAR6(P), PSD95(N-P3), TIP1(P), and KIAA0751(N3-J). There was weak expression (4-10% RFP positive) for DVL1(N), PAR6(N), and KIAA0751(P). CASK(P), DVL1(N-P), PAR(N-P), and TIP1(N-C) expression levels were 10-40% in Jurkat E6 cells. The DsRED plasmid was used as a RFP positive control for each set of transfections.

Fluorescent microscopy of Jurkat E6 was similar to the 293 HEK cells, the control DsRED construct was evenly diffused, varying from weak to very brightly fluorescence. The PRISM-RFP fusion protein staining patterns in Jurkat E6 are summarized as follows: CASK(P) some bright and diffuse and some punctated, DVL1(N-P) punctated and polarized and globular, DVL1(P) occasional diffuse, PAR6(N-P) polarized and globular and a few

compartmentalization in lymphocytes. Diminished expression of DVL1(N) and (P) and PAR6(N) and (P), as compared to DVL1(N-P) and PAR6(N-P) correlates with the loss of these domains. There also appears to be a hierarchy or an order to the successful localization of these molecules with the n-terminal domain being of first importance and the P (PDZ) domain playing a secondary role. In the case of TIP1(P) and TIP(N-C) the additional sequences on both the c- and n-terminal domains rescue lymphocytic expression.

7.4.4 References

Graham, F. and van der Eb, A., (1973). *Virology* 52:456.

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Potter, H. (1995). Recombinant DNA Methodology II. Academic Press, Inc. Chapter 31, pg. 467-484. Applications of Electroporation in Recombinant DNA Technology.

Reynaud et al. 2000, *J. Biol. Chem.* 275:33962-333968.

Suzuki et al, 1999, *Oncogene* 18:5967-72.

7.5 EXAMPLE 5: TIP1-RFP OVEREXPRESSION ENHANCES ANTI-CD95 INDUCED APOPTOSIS IN JURKAT T CELLS

This example shows the use of the assays and PRISM MATRIX described herein to identify a medically significant PDZ-PL interaction.

Human T cell lymphocyte virus type 1 (HTLV-1) is the etiologic agent of neoplasia within human peripheral blood T cells. The most important factor contributing to the initial stages of viral-mediated transformation of T cells after HTLV-1 infection is the viral oncoprotein Tax. Tax has been reported to bind to several proteins containing PDZ domains including TIP1 (Suzuki, T., et al., and Rousset et al., et al.).

Reviewing the PRISM MATRIX (similar to that provided in TABLE 2) we found that TIP1 binds to the CD95 and Tax. As described *infra*, we also determined that Tax competes for binding to TIP1 1000-fold better than CD95. Therefore we hypothesized TIP1 is a positive regulator of apoptosis and that in HTLV-1 infection, Tax blocks TIP1 binding to CD95 to promote an anti-apoptotic effect. We predicted that overexpression of TIP1-RFP would lower sensitivity to CD95-induced apoptosis and that TAX-1 (sometimes referred to herein as "TAX") can immortalize host cells by competing with CD95 for TIP-1 to block apoptosis.

A. TIP-1 Interaction with CD95 and TAX-1

5 As shown in the PRISM MATRIX, both the C-termini of Tax1 and of CD95 (Fas) bind to the PDZ domain of Tax1-interacting protein (Tax1-IP, TIP-1). Various concentrations of peptide ligands corresponding to the C-terminal 20 amino acids of Tax1 and CD95 were reacted with the Tax1-IP GST/PDZ fusion protein in a "G assay" format. These experiments revealed that the interaction of Tax1 with Tax1-IP is of higher affinity than the
10 interaction of CD95 with Tax1-IP. We therefore anticipated that a peptide corresponding to the C-terminal 8 amino acids of Tax1 would be able to block efficiently the binding of the 20 amino acid CD95 peptide to Tax1-IP. In addition, we anticipated that a peptide corresponding to the C-terminal 8 amino acids of CD95 would less efficiently block the binding of the 20 amino acid Tax1 peptide to Tax1-IP.

15 As shown in **Figures 6 and 7**, these predictions were correct: 100 uM of the truncated Tax1 peptide effectively blocked binding of the longer CD95 peptide to Tax1-IP, whereas 500 uM of the truncated CD95 peptide was required to block binding of the longer Tax1 peptide. Thus, information in the PRISM MATRIX, in combination with experiments investigating the affinities of different PDZ-ligand interactions, is useful in the design of
20 inhibitors of PDZ-ligand binding of different potencies.

Additional experiments showed that the CD95- TIP-1 interaction occurs in vivo. using biotinylated peptides corresponding to the C-terminal 20 residues of CD95 or TAX we were able to "pull-down" TIP-1-rfp from a cell lysate from transfected 293 HEK cells. This interaction was blocked in the presence of peptides having the sequence of the C-terminus (i.e.,
25 8 C-terminal residues) of TAX-1 or CD95.

B. Enhancement of Apoptosis

Jurkat E6 T cells were transfected with either 20 µg of the RFP, TIP1-RFP, or PAR6(N-P)-RFP. Twenty-four hours after transfection, cells were treated with 100 ng/ml anti-
30 human CD95 (Immunotech #1504, mouse monoclonal anti-human CD95 clone CH11) for 2hr. at 37°C. To analyze the levels of apoptosis in anti-CD95-treated cells we used the annexin V-FITC flow cytometer assay (Immunotech, Cat. No.2375). We followed the protocol supplied

by the manufacturer, except that propidium iodide was omitted because the RFP and propidium iodide emission spectra are overlapping using an Argon Laser (488 nm). To test for apoptosis with annexin V, approximately 5×10^5 cells were washed once with PBS and resuspended in 500 μ l of annexin V binding buffer and 5 μ l of annexin V-FITC solution was added. Cells were
5 incubated for 10 minutes on ice, in the dark. Each sample was then analyzed using flow cytometry.

Since the RFP emission spectra is broad, detectable in two of the three channels (FL-2 = 560-590 and FL-3 = 605-725), only the green channel (FL-1 = 505-545) is left for testing. We circumvented this by doing the experiment in two steps. First we set a forward
10 scatter, side scatter dot plot and then separated live from dead cells using a green viability dye (Sytox, Molecular probes). Secondly, we used the live cell gate to test for apoptotic cells with annexin V-FITC. Since many cells are already dead from electroporation and each construct can vary in percentage of cell death, we only compared live RFP positive cells with live RFP negative cells for annexin V. Additionally there is little cell death since the cells are only
15 treated with anti-CD95 for 2hrs. Viable cells were analyzed using a FL-1 by FL-3 two-color dot plot, and showed 4 distinct populations: viable transfected cells expressing RFP only, viable non-transfected cells, negative for RFP and annexin V-FITC, apoptotic non-transfected cells positive for annexin V-FITC, and apoptotic transfected cells double positive for RFP and annexin V-FITC [Figure 5A].

20 The results showed that cells expressing TIP1-RFP have a 30% increase in apoptosis versus the TIP1-RFP negative cells [Figure 5B]. In contrast, cells expressing either RFP control or PAR6(N-P)-RFP showed little to no increase in apoptosis, when compared to untransfected cells. The finding that overexpression of TIP1-RFP results in an increased sensitivity to anti CD95-mediated apoptosis demonstrates the physiological relevance of the
25 TIP1/CD95 interaction in vivo. Furthermore, transfected cells expressing higher of TIP1-RFP were more sensitive to apoptosis than cells expressing lower levels of TIP1-RFP, confirming the pro-apoptotic effects of TIP1.

HTLV-1 infection results in adult T cell leukemia and tropical paraparesis (a model for Multiple Sclerosis). Restoration of apoptosis to HTLV-1 infected cells by interfering
30 with TAX-1 interaction with CD95 is a basis for the treatment of HTLV-1. Furthermore, modulation of TIP1-CD95 interactions may be a target for treatment of Multiple Sclerosis.

7.6 EXAMPLE 6: HPV E6 ONCOGENE AND PRISM

This example demonstrates the use of PL sequence motifs identified according to the invention in the prediction of biological function in an oncogenic virus.

Human papilloma virus (HPV) infection plays a role in development of cervical carcinoma. The oncoprotein responsible for this is the early gene E6 from strains 16, 18 and 31. E6 associates with p53 and shunts this tumor suppressor into the ubiquitin proteosomal pathway to affect transformation. Using the PL motifs disclosed herein, we noted that the E6 from oncogenic strains HPV16, 18 and 31 are PDZ ligands (PLs) with the carboxy-terminal E-T-Q-V/L. Similarly, the E6 of oncogenic strain HPV66 has the carboxy-terminus ESTV, which also matches the consensus PDZ binding motif.

We performed an expanded search of the HPV E6 proteins and discovered HPV70 E6 fits perfectly the described PDZ consensus ETQV, identical to HPV18 and 31. We can thus predict that HPV70 is likely oncogenic on the basis that E6 is a PDZ ligand. Other HPV strains with E6 proteins that are potential PLs (based on motifs) include 57 (RTSH), 2a (RTLH), 63 (LYII). Strains 77 (QSRQ) and 80 (GSIE) may also be PLs, although the motif match is less strong. This information is summarized in **TABLE 9**.

PDZ targets for carboxy-terminal peptides corresponding to the above described E6s are determined using the methods of the invention (e.g., the G assay and PRISM MATRIX assay). Inhibitors of the interaction of the PDZ and oncogenic E6 PLs are determined and identified using the methods of the invention and are useful for inhibition of E6-mediated transformation. Such inhibitors (e.g., small molecules, peptides or recombinant proteins) are administered to patients (e.g., by local application to the vaginal vault and the uterine cervix) to treat or prevent cervical carcinoma. Diagnostic assays for oncogenic HPV are carried out using the sequences corresponding to the HPV E6 PL to design polynucleotide (e.g., PCR) or antibody probes that distinguish E6 proteins that are PLs from those that are not PLs.

Table 9

HPV E6 C-TERMINAL SEQUENCES				
Strain	GI	C-TERMINAL E6 SEQUENCES	ONCOGENIC	PDZ LIGAND
61	9628574	TGPCTARWQP		NO
60	9628566	RQRSYCRNCIEK		NO
55	9628558	CWTSCMETILP		NO
50	9628550	CCRNCEYHEG	NO	NO
48	9628542	CRNCISHEGR	NO	NO
44	9628534	CFHCWTSCMETILP	NO	NO
38	9628526	GNWKGRCRHCKAIE	NO	NO
37	9628518	WKGLCRHCGSIG	NO	NO
56	9628582	TGSCLCWRHTSRQATESTV	YES	YES
5	9626033	RCMNCAPRCMENAPALRTSH	ND	YES?
2	9626032	HCMNCGSSCTATDPASRTLH	ND	YES?
1	4927719	WTGRCMSCCRSSRTRRETQL	YES	YES
6	60995	HSCCNRARQERLQRRRETQV	YES	YES
3	333048	GRWTGRCIACWRRPRTETQV	YES	YES
4		CAACWRSARRRRLQRRRETAL	YES	YES
5		CANCWQTRQRRLQRRNETQV	YES	YES
6		CSECWRPTRRPRLQRRRVETQV	YES	YES
7		CAVCWRPARRRRLQRRRQETQV	YES	YES
8	134508	RHCWTSNREDRRRIRRETQV	ND	YES
9	312092	VHKVRNKFKAACSLCRLYII	ND	YES
10	2911558	GHWRGSLHCWSRCMGQSRQ		?
11	2911565	QFHKVRRNWKGLCRHCGSIE		?
21	9628462	WKGICRLCKHFQ		NO
11	333026	WKGRCCLHCWTTMEDLLP	NO	NO
36	9628510	WKGICRQCKHFYNDW	NO	NO
29	9628502	WRGSCLYCWSRCMGQSPR	NO	NO
28	9628494	CQYCWLRCVTRIPQ	NO	NO
24	9628486	KVRRGWKGLCRQCKQI	NO	NO
22	9628470	VRDHWKGRCRHCKAIE	NO	NO
21	9628462	HKVRGSWKGICRLCKHFQ	NO	NO
20	9628454	FYLVRGSWKGICRLCKHFQ	NO	NO
4	9626597	TCYLIRGLWRGYCRNCIRKQ	ND	NO
54	1017782	RRFHCVRGYWKGRCLHCWKP		
5B	9626498	KVRNAWKGICRQCKHFYHDW		

74	1491796	NTWKGRCFHCWTTCMENILP			
75	2911544	EFHKVRNRWKGVCRRHCRVIE			
76	2911544	EFHKVRNRWKGVCRRHCRVIE			
47	9627136	KVRNAWKGVCRQCKHFYNDW		ND	NO
65	9626613	ACYLIRGLWRGYCRNCIRKQ			

	1990-1991	1991-1992	1992-1993	1993-1994	1994-1995	1995-1996	1996-1997	1997-1998	1998-1999	1999-2000	2000-2001	2001-2002	2002-2003	2003-2004	2004-2005	2005-2006	2006-2007	2007-2008	2008-2009	2009-2010	2010-2011	2011-2012	2012-2013	2013-2014	2014-2015	2015-2016	2016-2017	2017-2018	2018-2019	2019-2020	2020-2021	2021-2022	2022-2023	2023-2024	2024-2025	2025-2026	2026-2027	2027-2028	2028-2029	2029-2030	2030-2031	2031-2032	2032-2033	2033-2034	2034-2035	2035-2036	2036-2037	2037-2038	2038-2039	2039-2040	2040-2041	2041-2042	2042-2043	2043-2044	2044-2045	2045-2046	2046-2047	2047-2048	2048-2049	2049-2050	2050-2051	2051-2052	2052-2053	2053-2054	2054-2055	2055-2056	2056-2057	2057-2058	2058-2059	2059-2060	2060-2061	2061-2062	2062-2063	2063-2064	2064-2065	2065-2066	2066-2067	2067-2068	2068-2069	2069-2070	2070-2071	2071-2072	2072-2073	2073-2074	2074-2075	2075-2076	2076-2077	2077-2078	2078-2079	2079-2080	2080-2081	2081-2082	2082-2083	2083-2084	2084-2085	2085-2086	2086-2087	2087-2088	2088-2089	2089-2090	2090-2091	2091-2092	2092-2093	2093-2094	2094-2095	2095-2096	2096-2097	2097-2098	2098-2099	2099-2100	2100-2101	2101-2102	2102-2103	2103-2104	2104-2105	2105-2106	2106-2107	2107-2108	2108-2109	2109-2110	2110-2111	2111-2112	2112-2113	2113-2114	2114-2115	2115-2116	2116-2117	2117-2118	2118-2119	2119-2120	2120-2121	2121-2122	2122-2123	2123-2124	2124-2125	2125-2126	2126-2127	2127-2128	2128-2129	2129-2130	2130-2131	2131-2132	2132-2133	2133-2134	2134-2135	2135-2136	2136-2137	2137-2138	2138-2139	2139-2140	2140-2141	2141-2142	2142-2143	2143-2144	2144-2145	2145-2146	2146-2147	2147-2148	2148-2149	2149-2150	2150-2151	2151-2152	2152-2153	2153-2154	2154-2155	2155-2156	2156-2157	2157-2158	2158-2159	2159-2160	2160-2161	2161-2162	2162-2163	2163-2164	2164-2165	2165-2166	2166-2167	2167-2168	2168-2169	2169-2170	2170-2171	2171-2172	2172-2173	2173-2174	2174-2175	2175-2176	2176-2177	2177-2178	2178-2179	2179-2180	2180-2181	2181-2182	2182-2183	2183-2184	2184-2185	2185-2186	2186-2187	2187-2188	2188-2189	2189-2190	2190-2191	2191-2192	2192-2193	2193-2194	2194-2195	2195-2196	2196-2197	2197-2198	2198-2199	2199-2200	2200-2201	2201-2202	2202-2203	2203-2204	2204-2205	2205-2206	2206-2207	2207-2208	2208-2209	2209-2210	2210-2211	2211-2212	2212-2213	2213-2214	2214-2215	2215-2216	2216-2217	2217-2218	2218-2219	2219-2220	2220-2221	2221-2222	2222-2223	2223-2224	2224-2225	2225-2226	2226-2227	2227-2228	2228-2229	2229-2230	2230-2231	2231-2232	2232-2233	2233-2234	2234-2235	2235-2236	2236-2237	2237-2238	2238-2239	2239-2240	2240-2241	2241-2242	2242-2243	2243-2244	2244-2245	2245-2246	2246-2247	2247-2248	2248-2249	2249-2250	2250-2251	2251-2252	2252-2253	2253-2254	2254-2255	2255-2256	2256-2257	2257-2258	2258-2259	2259-2260	2260-2261	2261-2262	2262-2263	2263-2264	2264-2265	2265-2266	2266-2267	2267-2268	2268-2269	2269-2270	2270-2271	2271-2272	2272-2273	2273-2274	2274-2275	2275-2276	2276-2277	2277-2278	2278-2279	2279-2280	2280-2281	2281
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7.7 EXAMPLE 7: DESIGN OF SMALL MOLECULE INHIBITORS OF PDZ-LIGAND INTERACTIONS, AND THEIR CLINICAL APPLICATION

As shown in **TABLE 2**, both the C-termini of Dock2 and of BLR-1 bind to the PDZ domain of KIAA0807. This example describes the design of small molecule (<600 molecular weight) inhibitors of these interactions. As described *supra*, chemical entities resembling the C-termini of PDZ ligands that bind to a given PDZ domain (including peptides and mimetics) are effective inhibitors of ligand binding to that PDZ domain. We synthesized and purified 8 amino acid peptides corresponding to the C-termini of Dock2 and BLR-1, for use as inhibitors of KIAA0807-ligand binding. In addition, we synthesized and purified small molecules corresponding to the C-terminal four amino acids of Dock2 and BLR-1, acetylated at the N-terminus. Inhibition of KIAA0807-ligand binding by peptide (8 amino acid) and small molecule (4 amino acid) inhibitors are shown in **Figures 8 and 9**.

In **Figure 8**, the bars on the left hand side of the figure show that increasing concentrations of the peptide inhibitor (the C-terminal 8 amino acids of BLR-1) are somewhat effective at blocking binding of 1 uM of the biotinylated C-terminal 20 amino acids of BLR-1 to KIAA0807 GST/PDZ fusion protein. The bars of the right hand side of the figure show that increasing concentrations of the small molecule inhibitor (Acetyl-LTTF) are equally or more effective. In **Figure 9**, the bars on the left hand side of the figure show that increasing concentrations of the peptide inhibitor (the C-terminal 8 amino acids of Dock2) are somewhat effective at blocking binding of the 1 uM of the biotinylated C-terminal 20 amino acids of Dock2 to KIAA0807 GST/PDZ fusion protein. The bars on the right hand side of the figure show that increasing concentrations of the small molecule inhibitor (Acetyl-STD1) are equally or more effective. Thus, a general route to producing a small molecule inhibitor of a PDZ-ligand interaction is to synthesize a molecule corresponding to the C-terminal four amino acids of the involved ligand, acetylated at the N-terminus. This compound can subsequently be altered by art known means (e.g., changing its covalent composition to optimize pharmacokinetic properties without grossly altering its molecular structure, especially the molecular structure of the most C-terminal protein).

7.8 **EXAMPLE 8:** INHIBITION OF INDIVIDUAL INTERACTIONS FOUND IN
THE PRISM MATRIX

As shown in the PRISM MATRIX (TABLE 2), the C-terminus of Clasp-4 binds to the PDZ domain of KIAA440, and the C-terminus of DNAM-1 binds to the PDZ domain of WWP3. As disclosed herein, it was anticipated that (i) a molecule resembling (i.e., structurally similar to) the C-terminus of Clasp-4 would inhibit the Clasp-4/KIAA440 interaction and (ii) a molecule resembling the C-terminus of DNAM-1 would inhibit the DNAM-1/WWP3 interaction.

Peptides corresponding to the C-terminal 8 amino acids of Clasp-4 and DNAM-1 were synthesized and purified. These peptides were then tested for their abilities to inhibit the corresponding interactions. Inhibitor peptide concentrations in the range of 20 – 200 uM were effective (i.e., at least 50% inhibition) at blocking the interactions of labeled peptides corresponding to the C-termini of Clasp-4 and DNAM-1 with KIAA440 and WWP3, respectively (as measured in a "G assay" format). Thus, as described hereinabove, for each interaction identified in the PRISM MATRIX, it is possible to design or identify (e.g., by library screening) an effective inhibitor of that interaction by synthesizing a molecule (e.g., a peptide, peptide mimetic, or structurally similar small organic molecule) resembling the C-terminus of the PL involved in the interaction.

The present invention is not to be limited in scope by the exemplified embodiments which are intended as illustrations of single aspects of the invention and any sequences which are functionally equivalent are within the scope of the invention. Indeed, various modifications of the invention in addition to those shown and described herein will become apparent to those skilled in the art from the foregoing description and accompanying drawings. Such modifications are intended to fall within the scope of the appended claims.

All publications cited herein are incorporated by reference in their entirety and for all purposes.

WHAT IS CLAIMED IS

1. A method of modulating a biological function of an endothelial cell or hematopoietic cell, comprising introducing into the cell an agent that inhibits binding of a PDZ protein and a PL protein in the cell, thereby modulating the biological function.
2. The method of claim 1, wherein the PL is selected from the group consisting of CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, , CD44, CD38, CD3n, LPAP, CD46, CDw128B, DOCK2, PAG, CD34, and BLR-1.
3. The method of claim 2, wherein the PDZ is selected from the group consisting of MPP1, K303, K807, DLG1, PSD95, NeDLG, IP43, LDP, LIM, K545, TIP1, PTN-4, CBP, AF6, PDZK1, DLG5, Syntenin, WWP3, and K561.
4. The method of claim 1, wherein
- a) the PDZ protein is MPP1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y/I-X-V;
 - b) the PDZ protein is LIMK1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V;
 - c) the PDZ protein is K303 and the PL protein has a carboxy-terminal amino acid motif X-S-X-V;
 - d) the PDZ protein is K807 and the PL protein has a carboxy-terminal amino acid motif X1-S/T-X2-V/I/L/F;
 - e) the PDZ protein is DLG1, PSD95, or NeDLG and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y/A/E-X-V/I/L;
 - f) the PDZ protein is SNTa1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-D/Y-V/I/L;
 - g) the PDZ protein is DVL1 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V;
 - h) the PDZ protein is LDP and the PL protein has a carboxy-terminal amino acid motif X-A/S-X2-V/I;
 - i) the PDZ protein is LIM and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-A/V;
 - j) the PDZ protein is K561 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-V/I/L/F;

k) the PDZ protein is K545 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y-M-A/S/V;

l) the PDZ protein is TAX-IP2 and the PL protein has a carboxy-terminal amino acid motif X-S-D/E-V;

5 m) the PDZ protein is MPP2 and the PL protein has a carboxy-terminal amino acid motif X-S/T/Y-X-A/V/I;

n) the PDZ protein is TIP-1 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-V/I/L;

10 o) the PDZ protein is PTN-4 and the PL protein has a carboxy-terminal amino acid motif X1-S/T-X-V/F;

p) the PDZ protein is prIL16 and the PL protein has a carboxy-terminal amino acid motif D/E/K/R-V/I/L/F/Y-X-V;

q) the PDZ protein is CBP and the PL protein has a carboxy-terminal amino acid motif X-S/T-F/Y-V;

15 r) the PDZ protein is protein 41 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y/F-X-A/V/I/L;

s) the PDZ protein is AF6 and the PL protein has a carboxy-terminal amino acid motif X-A/S/T/Y-F/Y-V/I/L;

20 t) the PDZ protein is RGS12 and the PL protein has a carboxy-terminal amino acid motif X1-S/T/Y-X-V/F;

v) the PDZ protein is PDZK1 and the PL protein has a carboxy-terminal amino acid motif X-T-X-F;

w) the PDZ protein is DLG5 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X-V;

25 x) the PDZ protein is Synt and the PL protein has a carboxy-terminal amino acid motif X1-V/I/L-X2-V;

y) the PDZ protein is WWP3 and the PL protein has a carboxy-terminal amino acid motif X-S/T-X2-V; and,

30 z) the PDZ protein is TAX-IP40 and the PL protein has a carboxy-terminal amino acid motif X-Y-X-V;

where X is any amino acid, X1 is any amino acid, X2 is any amino acid.

5. The method of claim 2 wherein the agent is a peptide comprising a sequence of at least the carboxy-terminal two residues of the PL protein.

6. The method of claim 4 wherein the agent is a peptide comprising a sequence of at least the carboxy-terminal three residues of the PL protein.

7. The method of claim 2 wherein the agent is a small molecule or peptide mimetic of the carboxy-terminus of the PL protein.

8. The method of claim 4, wherein the cell is a T cell or a B cell.

9. A method of determining whether a test compound is an inhibitor of binding between a PDZ protein and a PL protein comprising:

a) contacting

i) a PDZ domain polypeptide having a sequence from the PDZ protein, and

ii) a PL peptide, wherein the PL peptide comprises a C-terminal sequence of a PL protein selected from the group consisting of CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, TAX, CD44, CD38, CD3 η , LPAP, CD46, CDw128B, DOCK2, PAG, CD34, and BLR-1 under conditions in which they form a complex

wherein said contacting is carried out in the presence and in the absence of a test compound;

b) detecting the formation of the complex in the presence and absence of the test compound

wherein less complex formation in the presence of the test compound than in the absence of the compound indicates that the test compound is an inhibitor of a PDZ protein -PL protein binding.

10. An inhibitor identified by the method of claim 9.

11. The inhibitor of claim 9 that is

(a) a peptide comprising a sequence that is from 3 to about 20 residues of a C-terminal sequence of a PL protein selected from CD105, VCAM1, CD95, Spectrin β , KV1.3, DNAM1, Neuroligin 3, TAX, CD44, CD38, CD3 η , LPAP, CD46, CDw128B, DOCK2, PAG, CD34, and BLR-1;

(b) a peptide mimetic of a peptide of section (a); or

(c) a small organic molecule with a molecular weight less than 1 kD.

12. A pharmaceutical composition comprising an inhibitor of claim 11.

13. A method for treating a disease characterized by leukocyte activation, comprising administering a therapeutically effective amount of an inhibitor of claim 11.

5 14. The method of claim 13 wherein the disease is characterized by an inflammatory or humoral immune response.

15. The method of claim 14 wherein the disease is an autoimmune disease.

10 16. A method of modulating a biological function of a cell, comprising introducing into the cell an antagonist that inhibits binding of a PDZ protein and a PL protein in the cell, wherein,

a) the PDZ protein is MPP1 (p55) and the PL is Spectrin β ;

b) the PDZ protein is K303 and the PL is Spectrin β ;

15 c) the PDZ protein is K807 and the PL VCAM1, Spectrin β , KV1.3, Neuroligin 3, CD38, CD3 η , LPAP, CD46 (form 1), CDw128B, DOCK2, PAG, CD34, or BLR-1;

d) the PDZ protein is DLG1 and the PL is Spectrin;

e) the PDZ protein is PSD95 and the PL is Spectrin β , CD34, or CD38;

20 f) the PDZ protein is NeDLG and the PL is Spectrin β or CD38;

g) the PDZ protein is TAX IP43 and the PL is Spectrin β or CD38;

h) the PDZ protein is LDP and the PL is CD38;

i) the PDZ protein is LIM and the PL is CD105;

j) the PDZ protein is K545 and the PL is CD105;

25 k) the PDZ protein is TIP1 and the PL is CD95, KV1.3, CD3 η , LPAP;

l) the PDZ protein is PTN-4 and the PL is Spectrin β ;

m) the PDZ protein is CBP and the PL is Spectrin β ;

n) the PDZ protein is AF6 and the PL is Spectrin β ;

o) the PDZ protein is PDZK1 and the PL is BLR-1;

30 p) the PDZ protein is DLG5 and the PL is Spectrin;

q) the PDZ protein is Syntenin and the PL is CD44;

r) the PDZ protein is WWP3 and the PL is VCAM1, Spectrin β , DNAM1,

Neurologin 3;

s) the PDZ protein is K561 and the PL is BLR-1.

17. The method of claim 16 wherein the cell is a hematopoietic cell.

5

ABSTRACT

MOLECULAR INTERACTIONS IN HEMATOPOIETIC CELLS

5 The invention provides reagents and methods for inhibiting or enhancing interactions between proteins in hematopoietic cells and other cells involved in the mediation of an immune response. Reagents and methods provided are useful for treatment of a variety of diseases and conditions mediated by immune system cells.

10 PA 3112092 v1

FIG. 1A

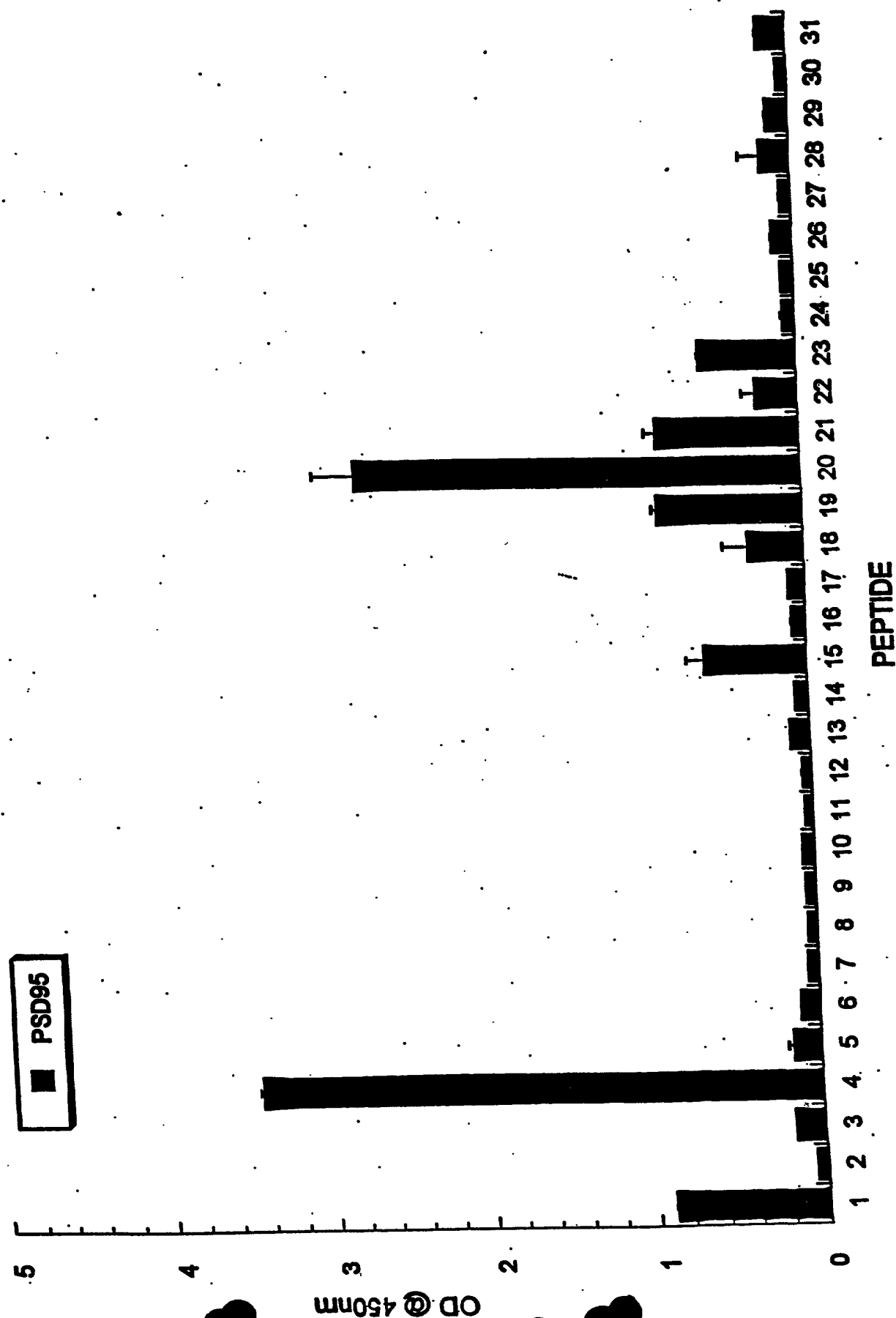


FIG. 1B

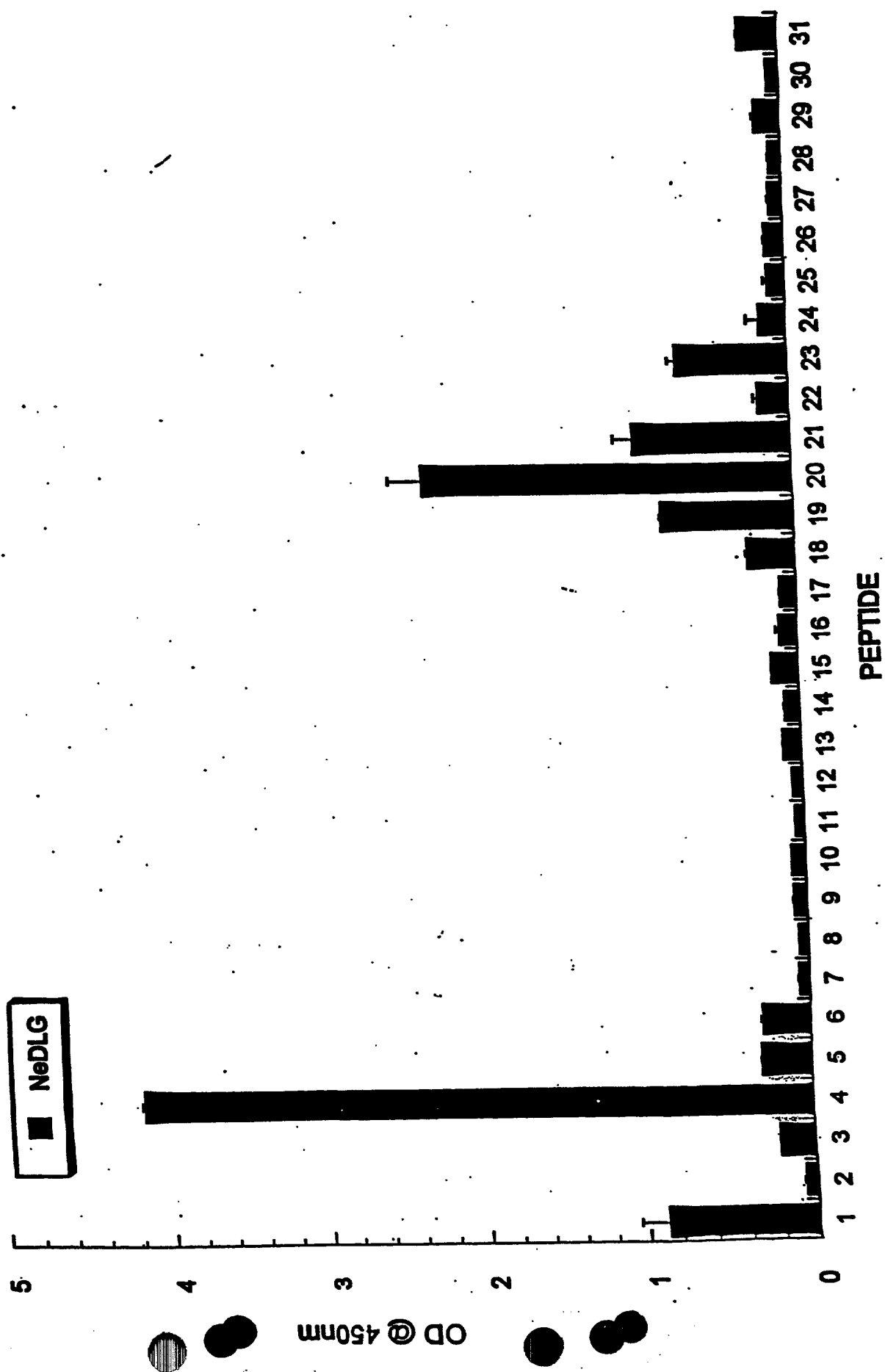


FIG. 1D

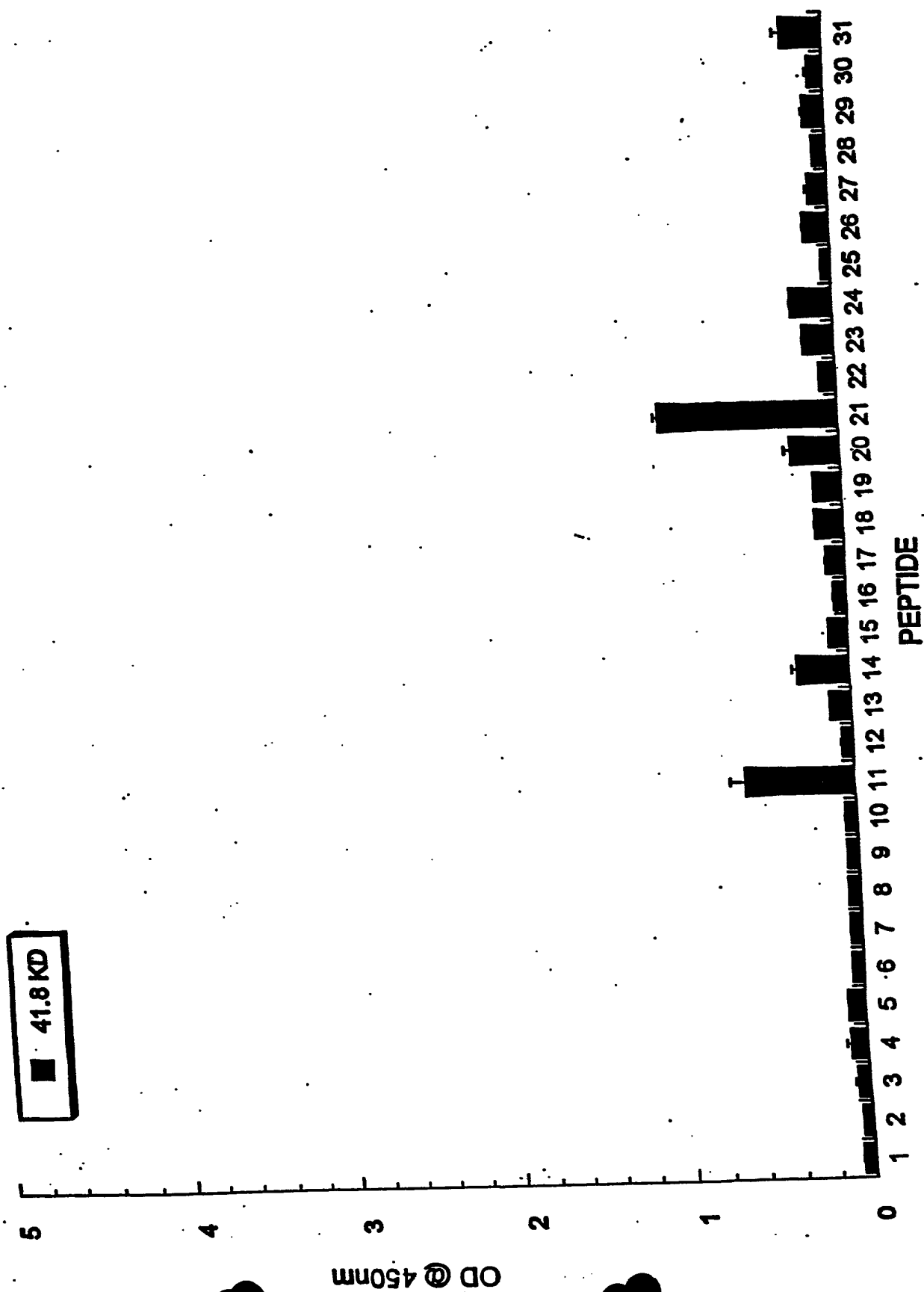


FIG. 2A

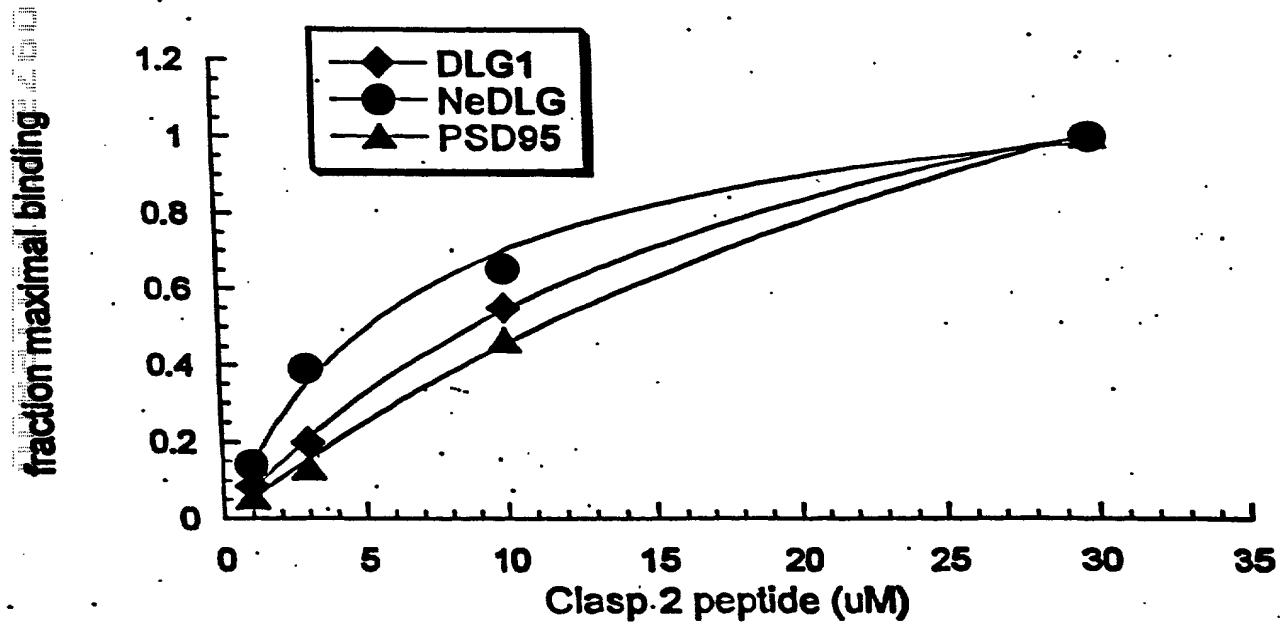
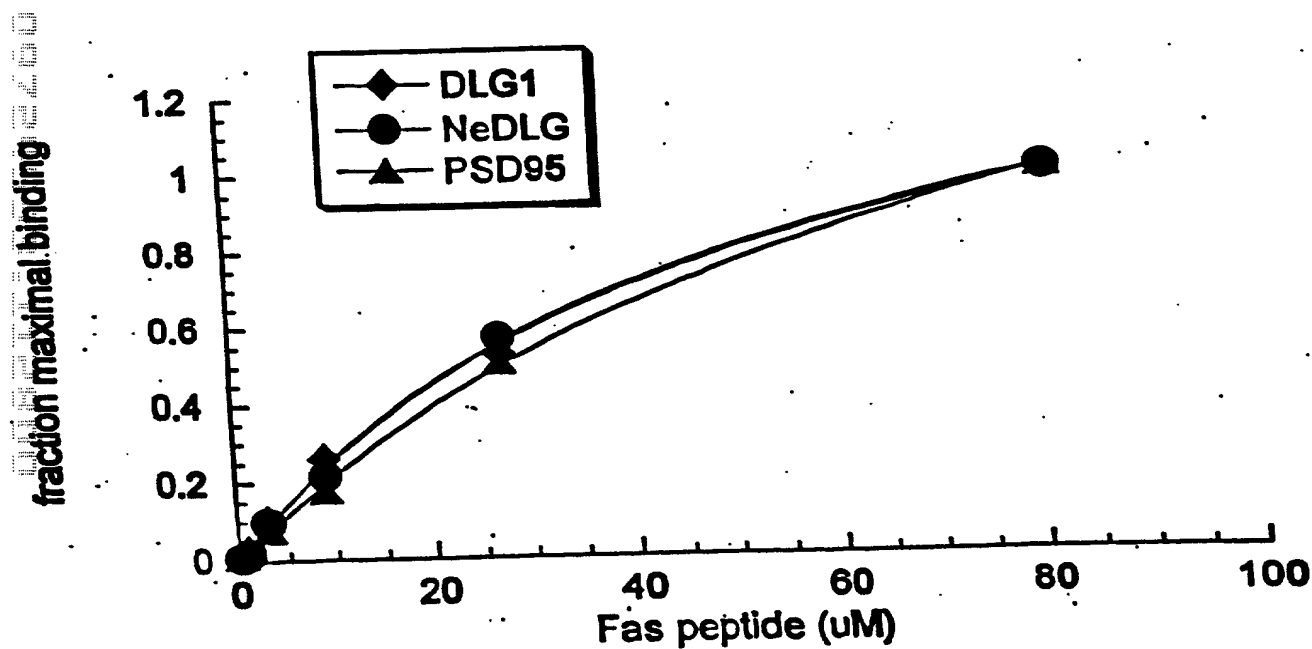
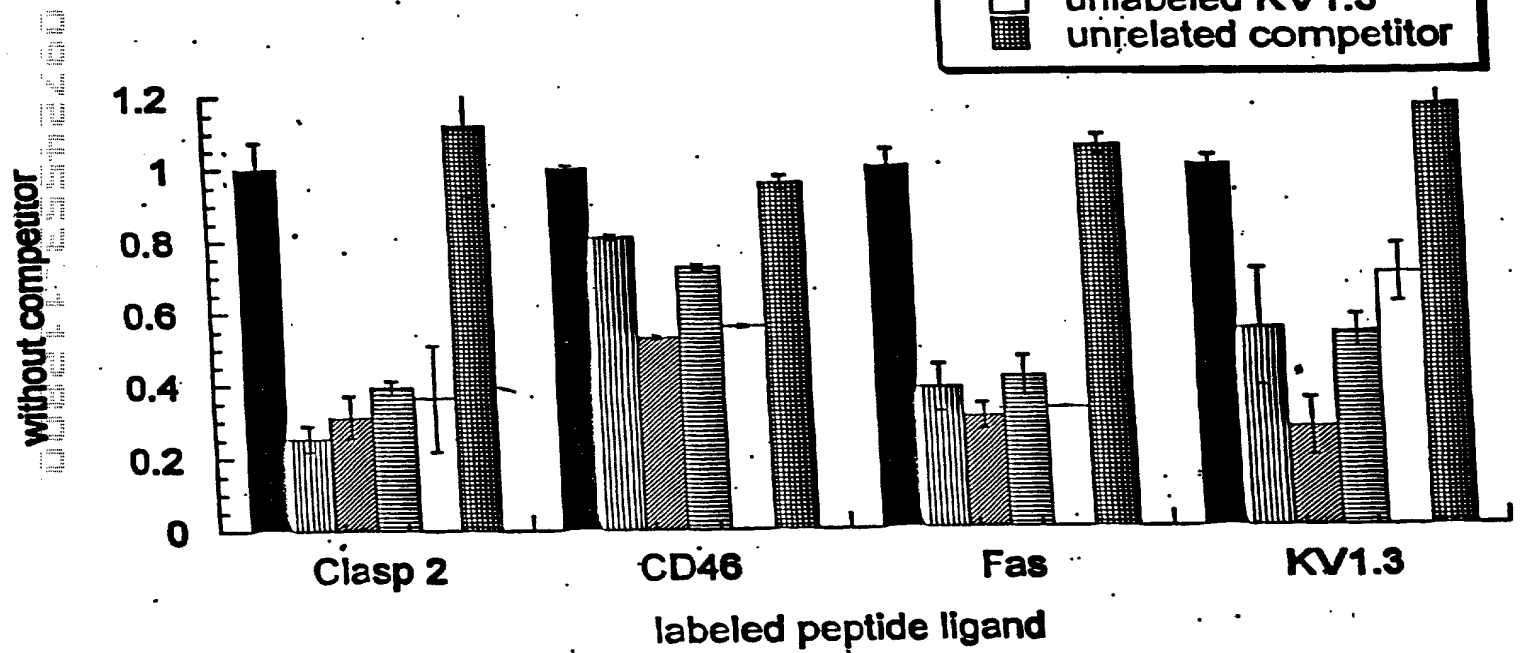


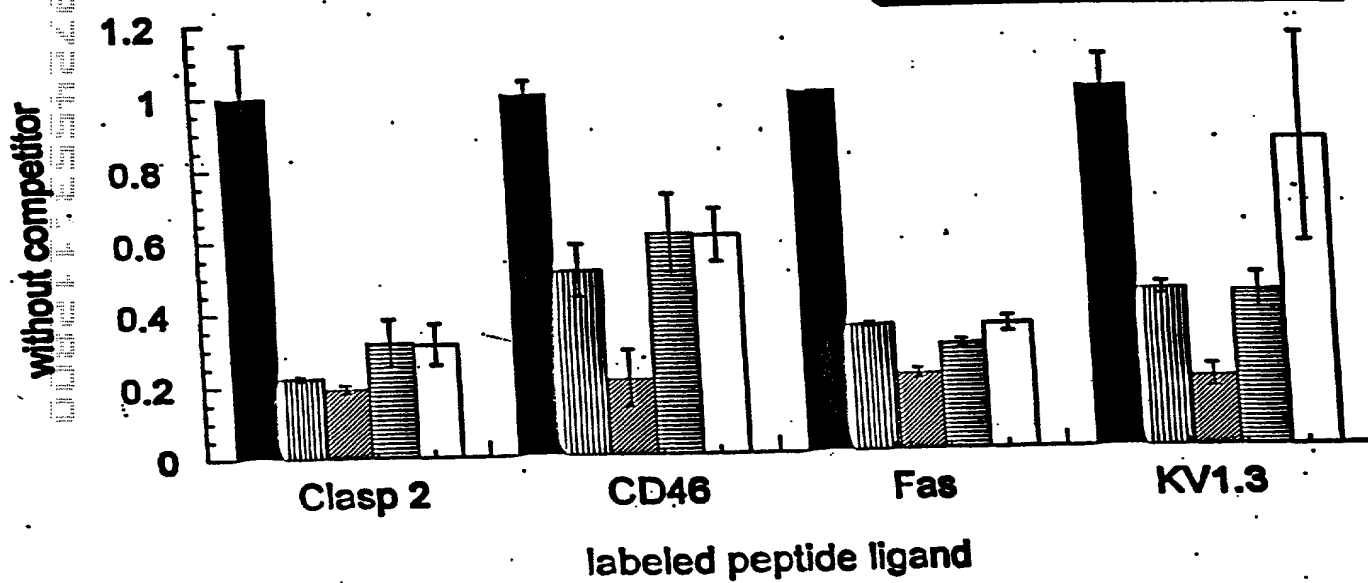
FIG. 2B



PDZ DLG1-GST Fusion Protein



PDZ PSD95-GST Fusion Protein



PDZ NeDLG-GST Fusion Protein

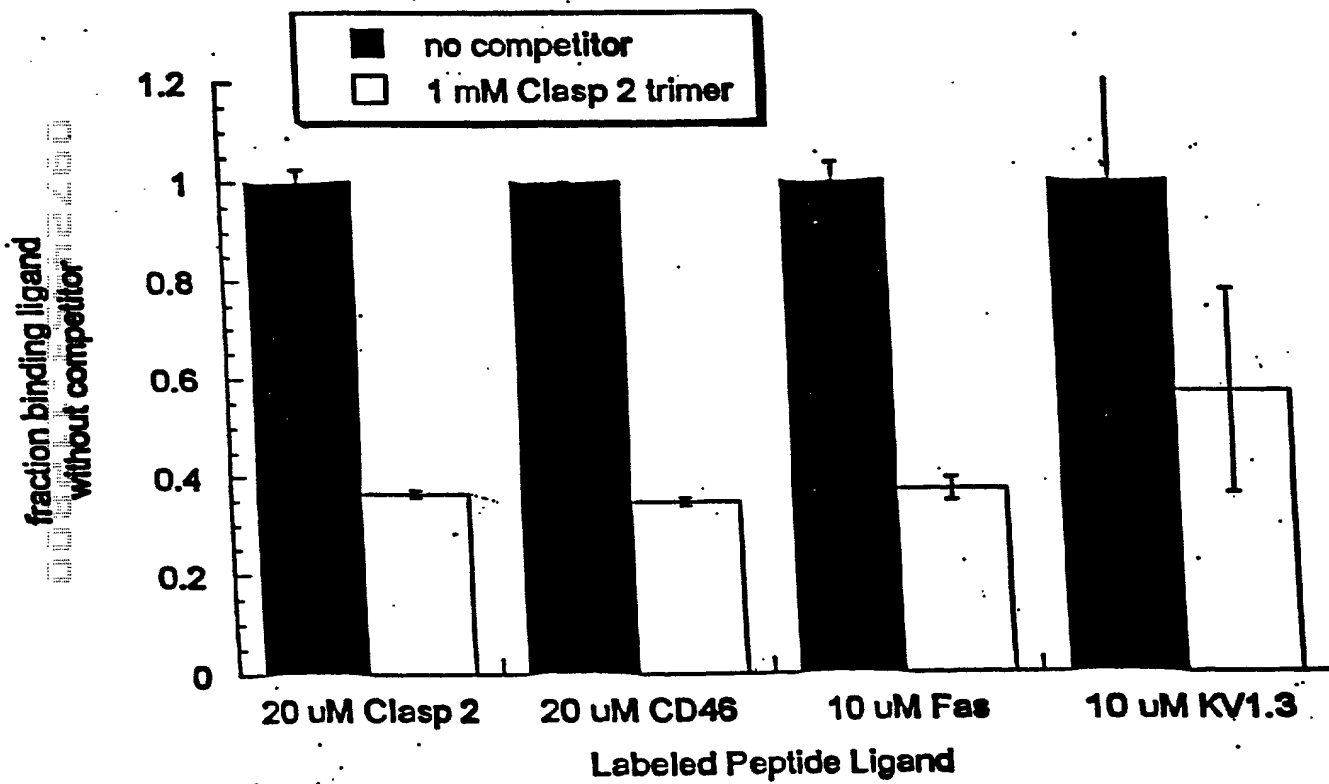


FIG. 3D

PDZ DLG1-GST Fusion Protein

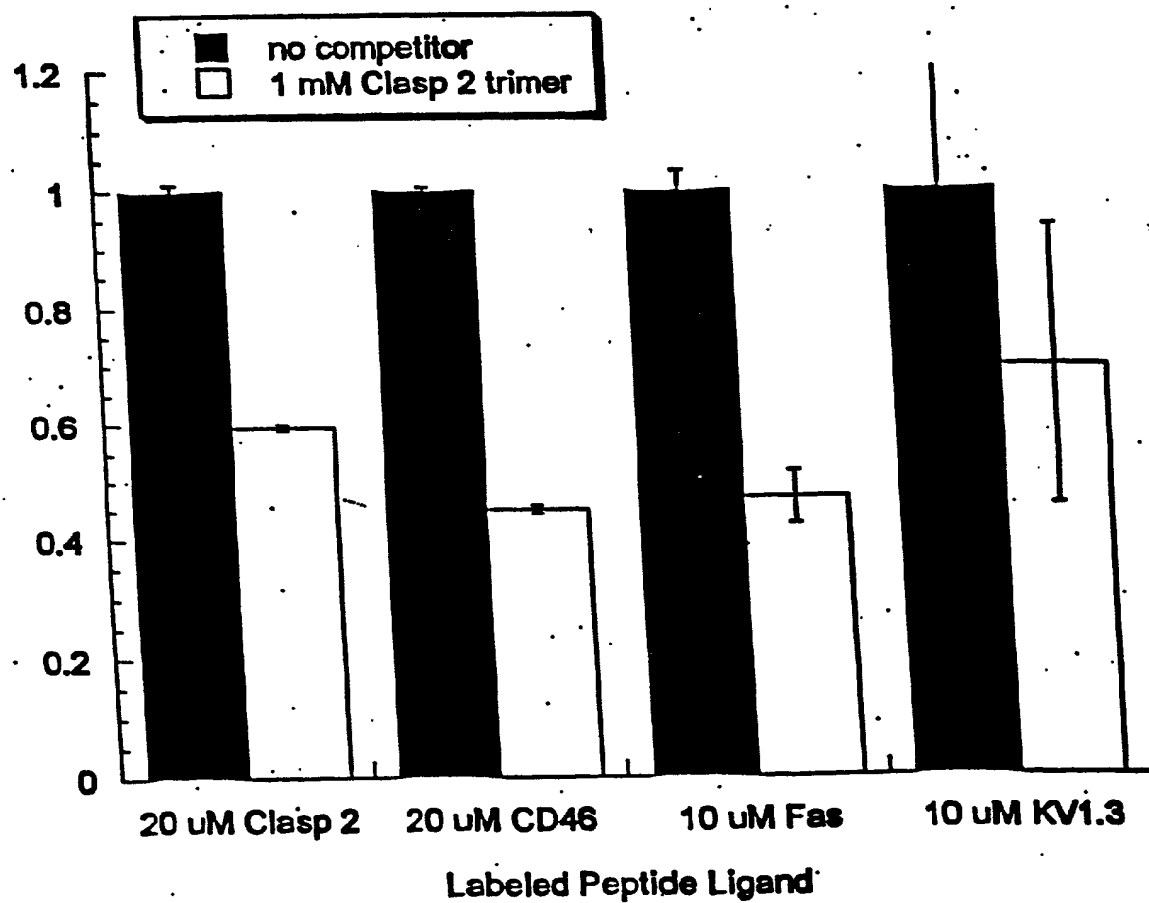


FIG. 3E

PDZ PSD95-GST Fusion Protein

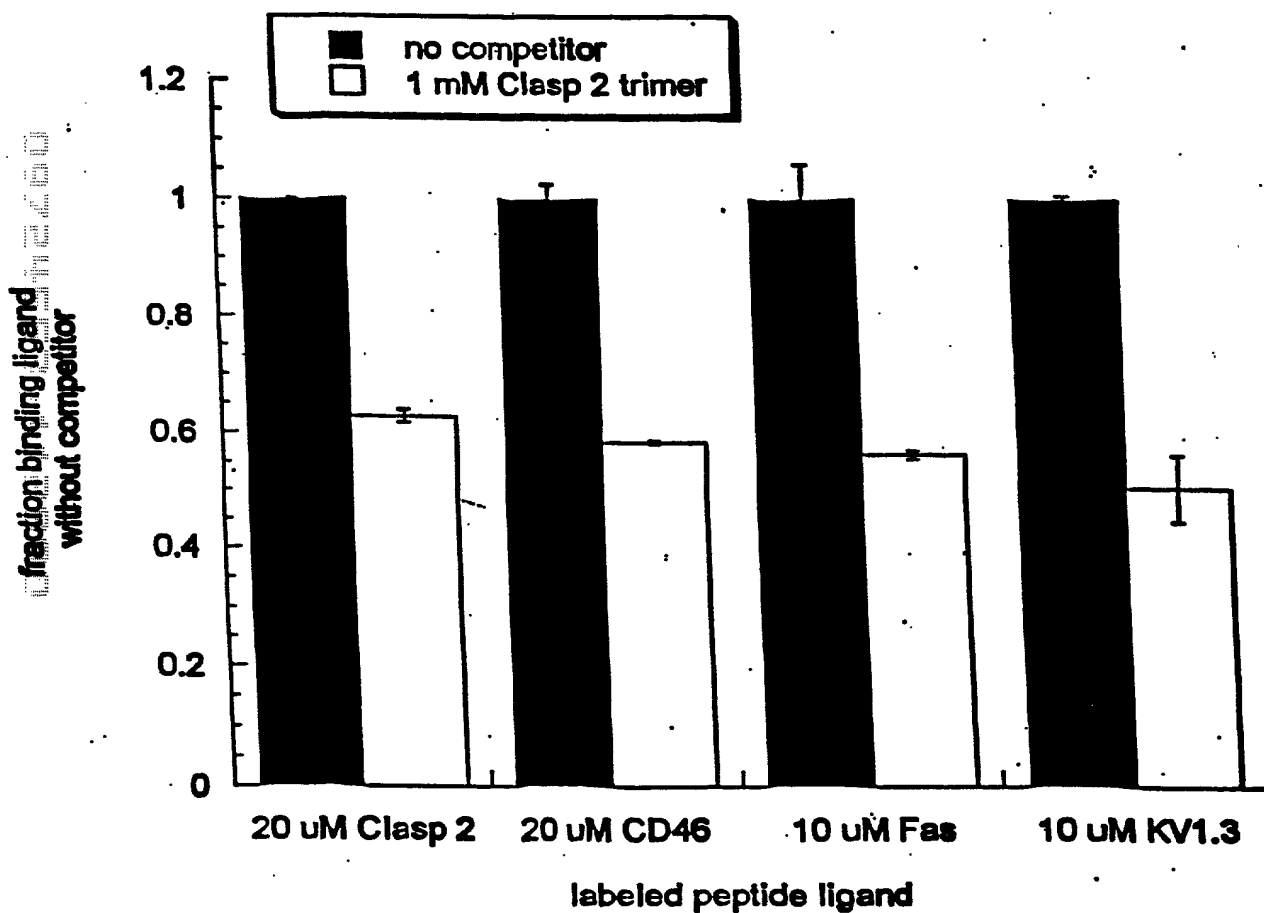
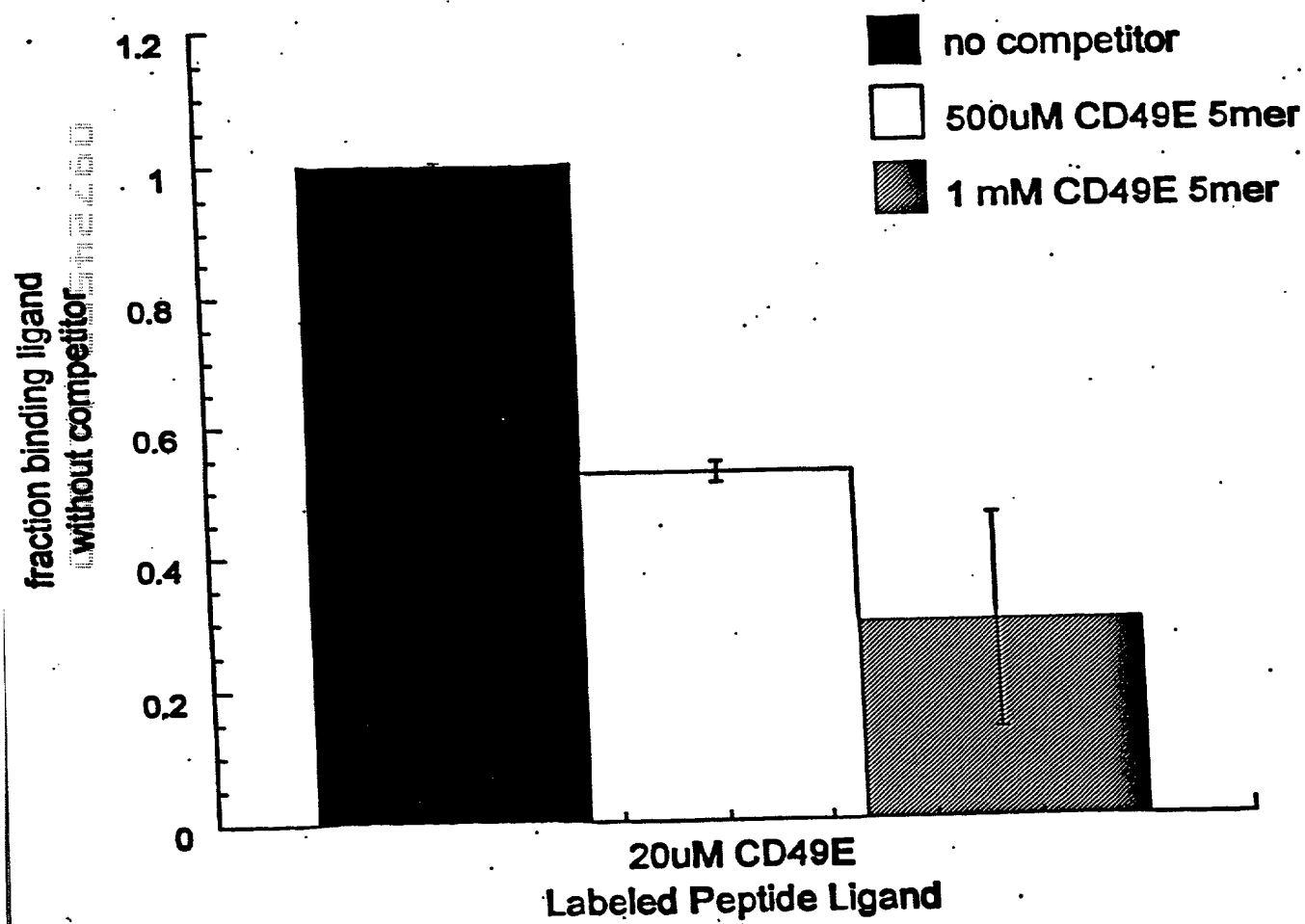
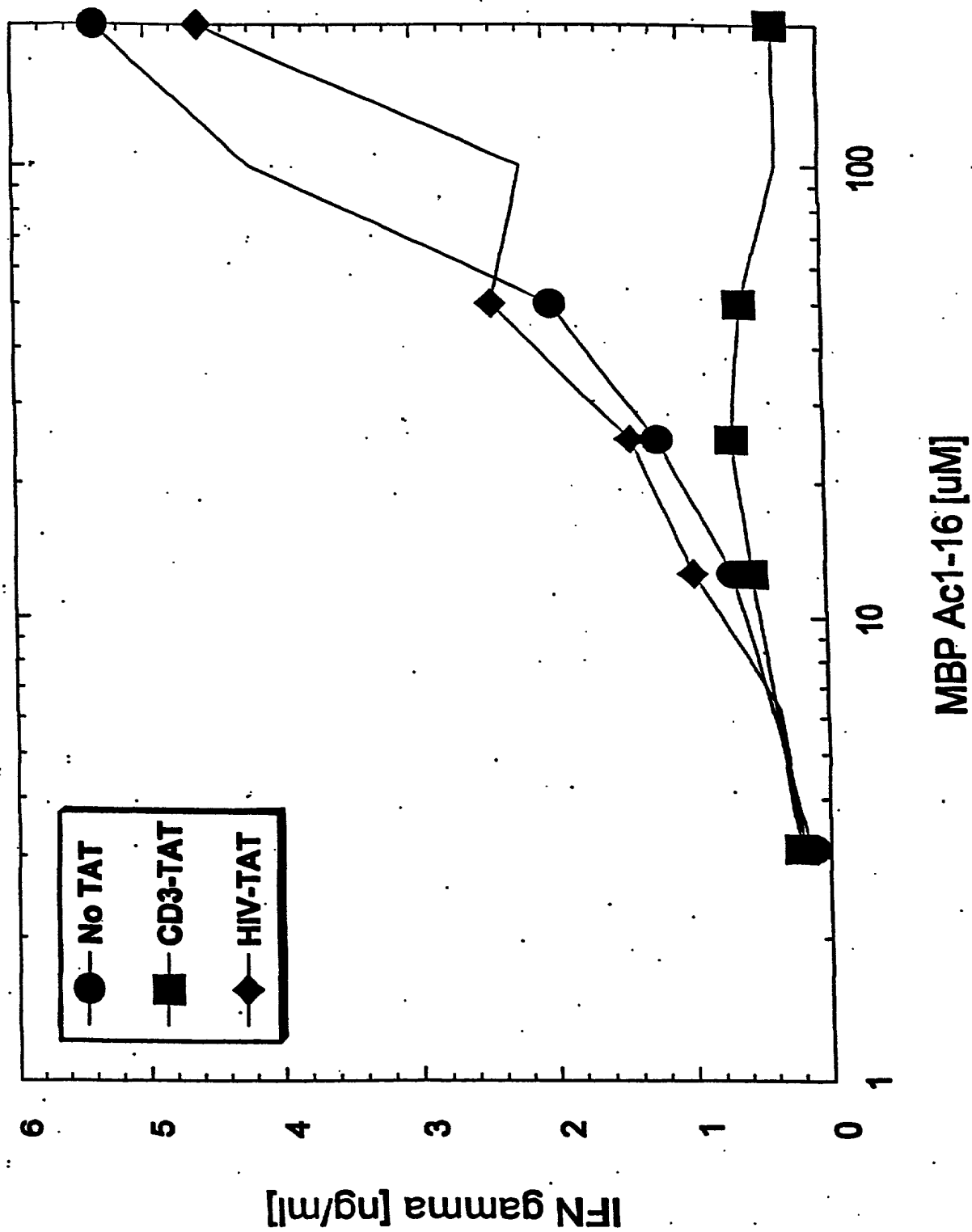


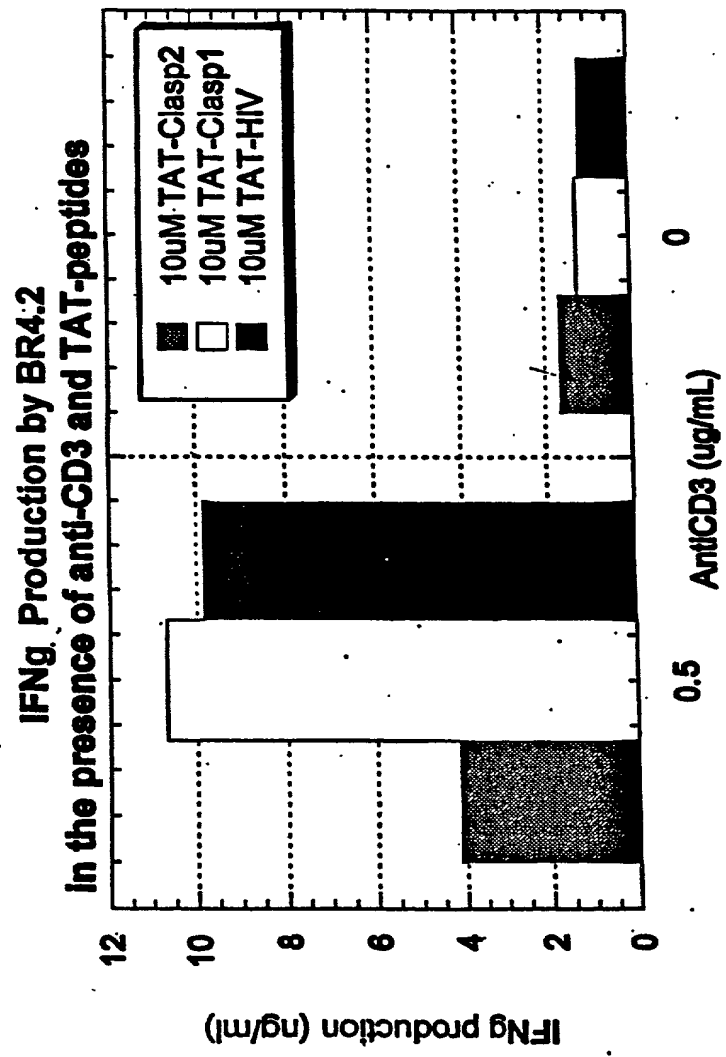
FIG. 3F

PDZ 41.8kD-GST Fusion Protein



IFN gamma response of
BR4.2 T cells stimulated with B10.BR:MBP Ac1-16 +/-
HIV-TAT or CD3-TAT peptides @ 10uM



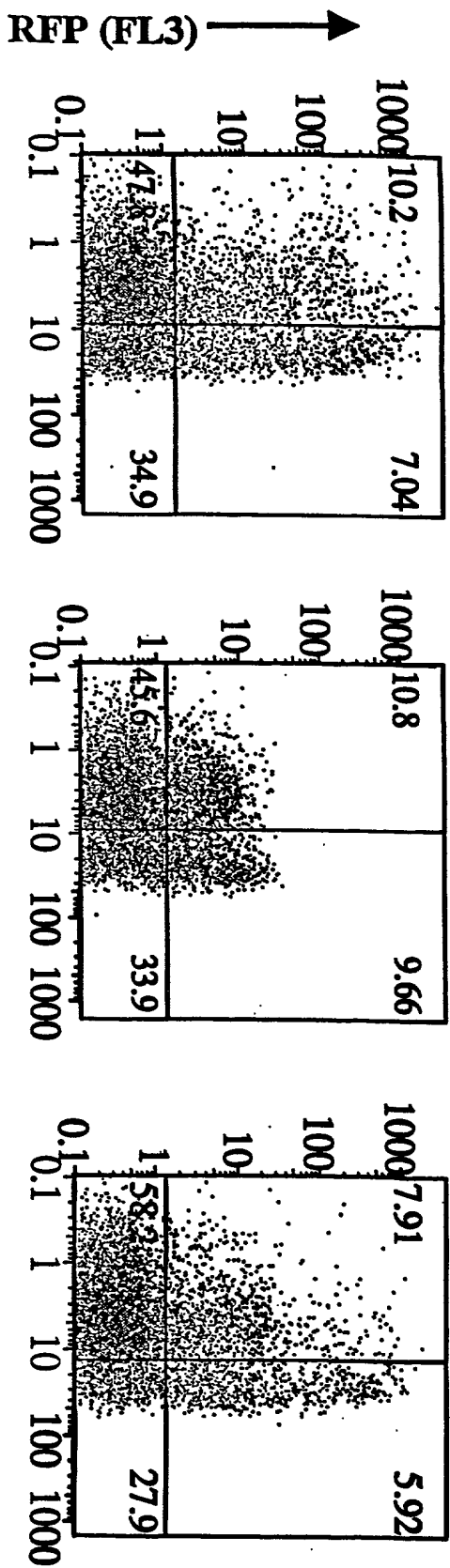


A.

DSRED

PAR6 (N-P)-RFP

TIP-1 (N-C)-RFP



B.

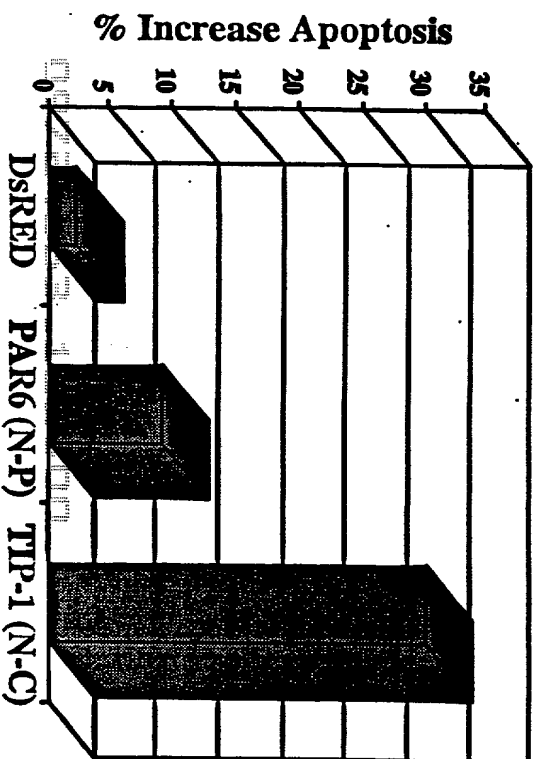


FIGURE 5

Fig. 6

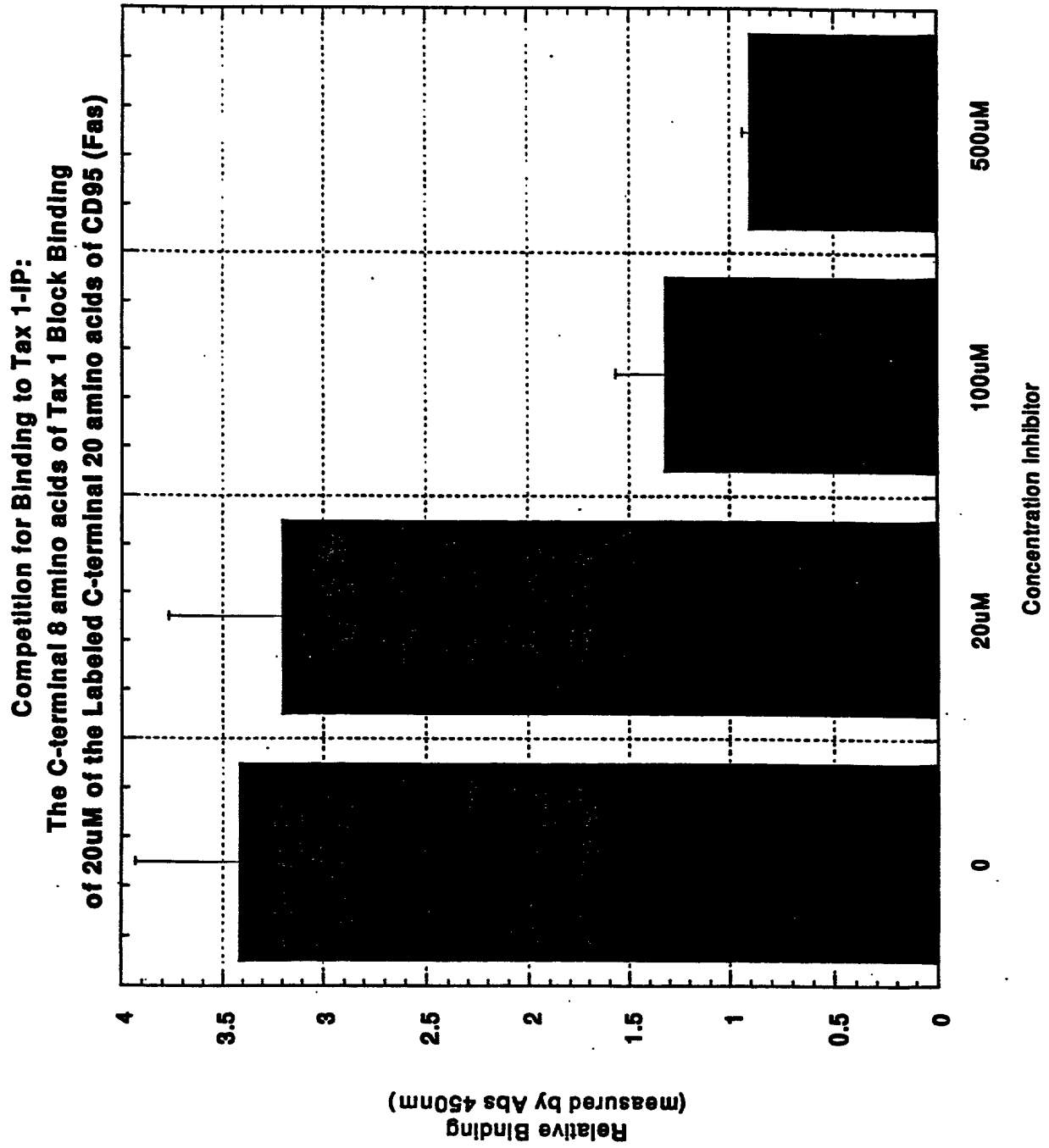
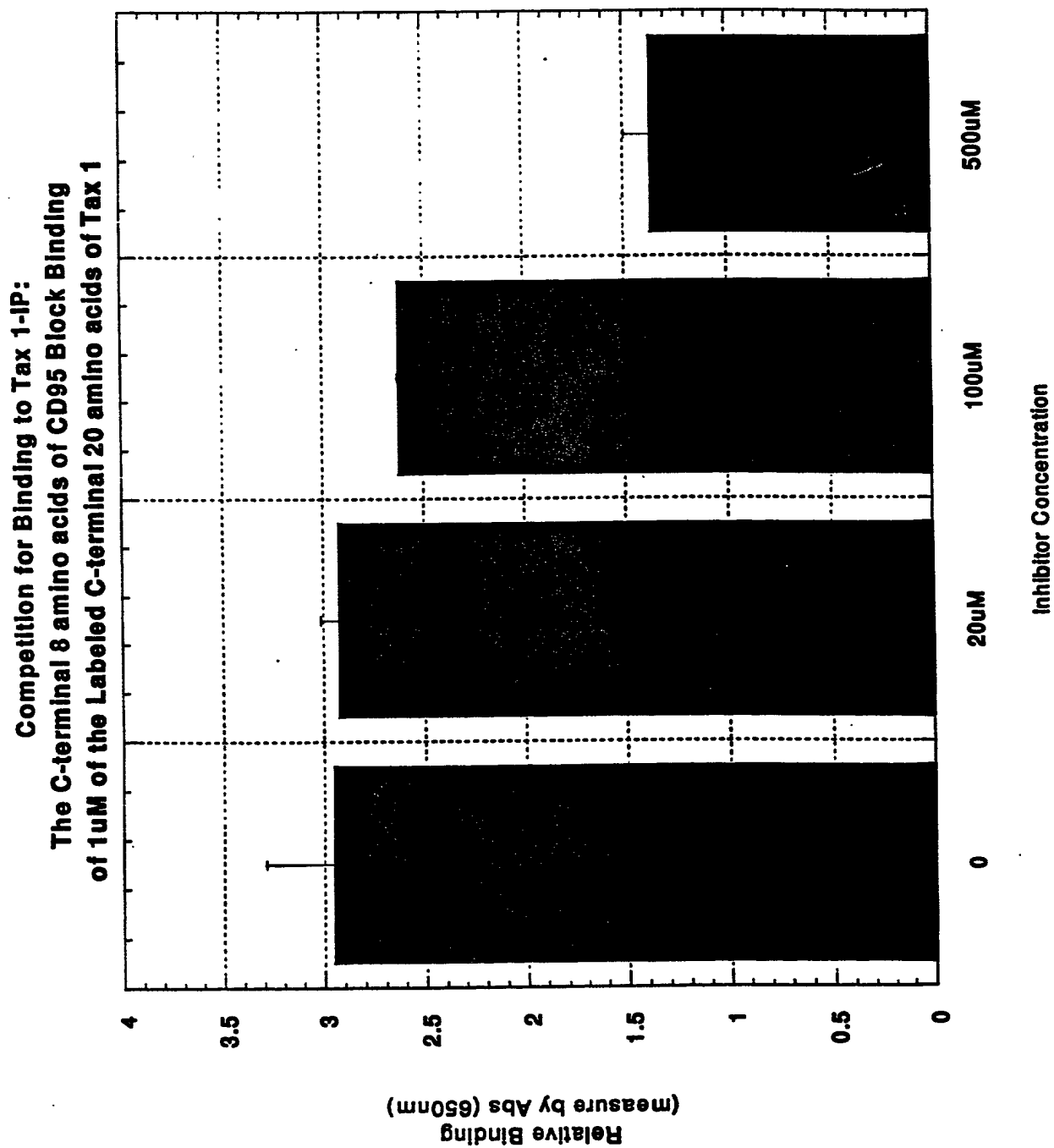


Fig. 7



**Inhibition of Binding of 1 μ M Biotin Labeled Peptide
Corresponding to the C-terminal 20 amino acids of BLR-1
to KIAA0807:GST Fusion Protein by Peptide
and Small Molecule Antagonists**

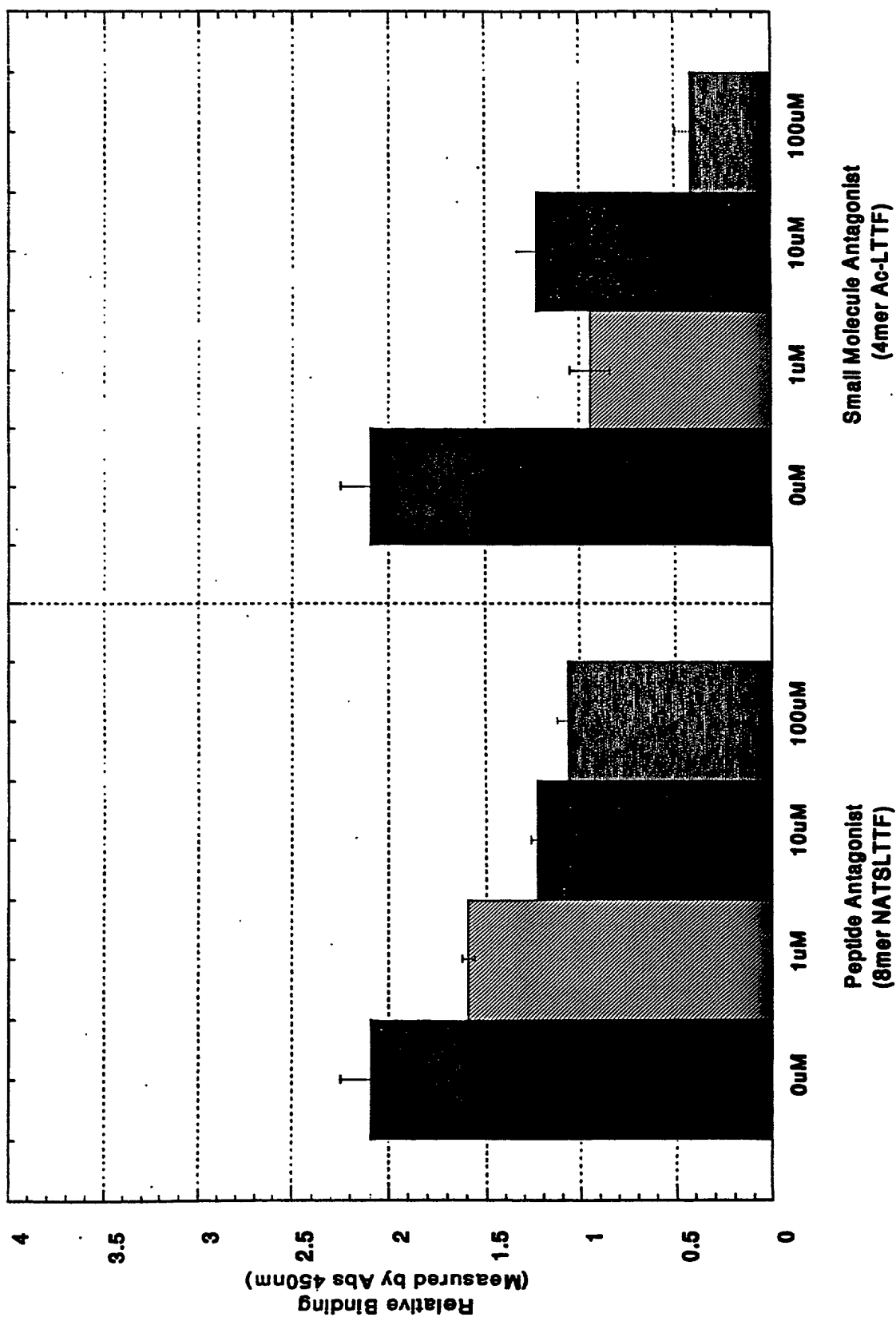
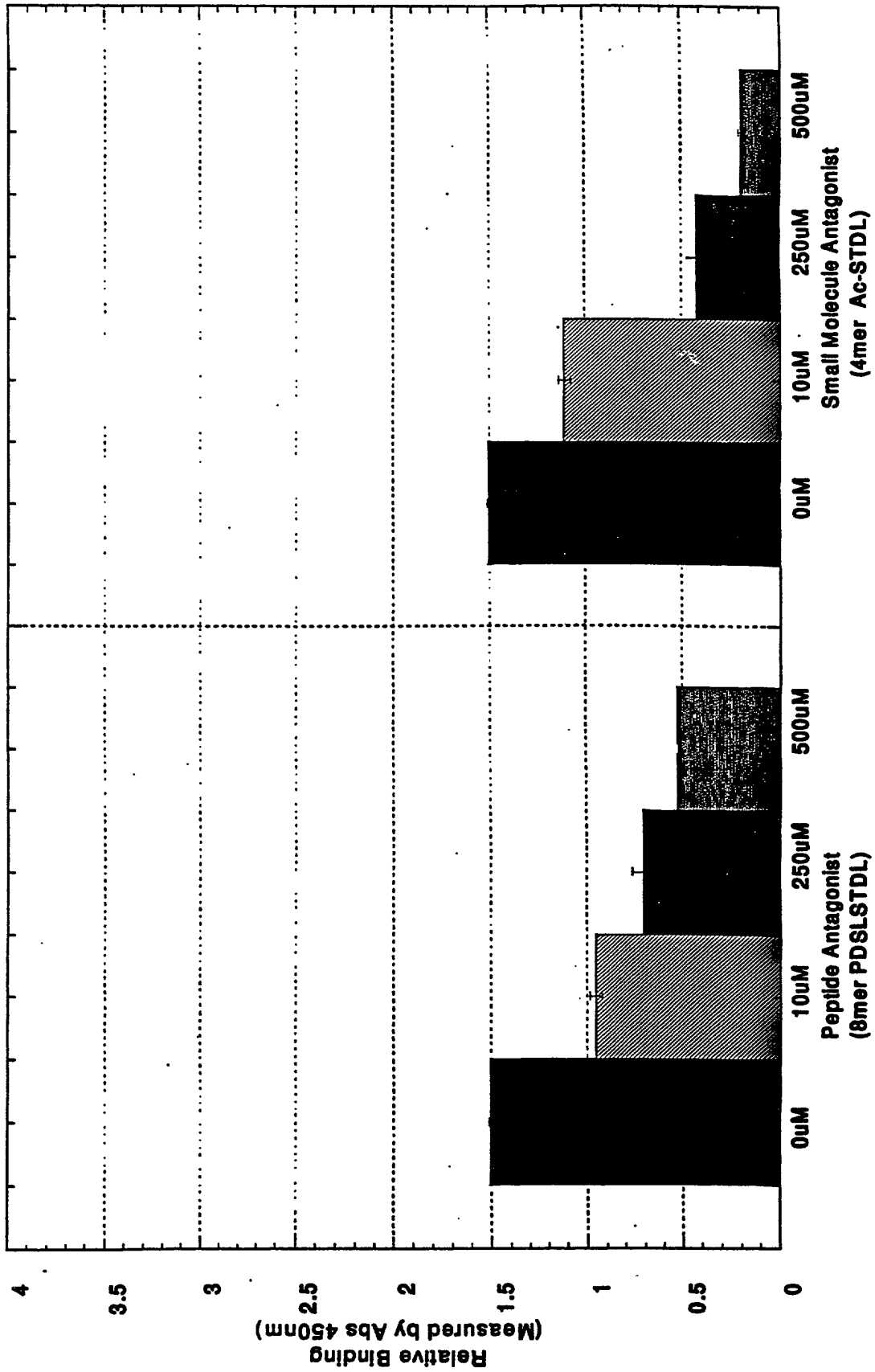


Fig. 8

Fig. 9

**Inhibition of Binding of 10uM Biotin Labeled Peptide
Corresponding to the C-terminal 20 amino acids of DOCK2
to KIAA0807:GST Fusion Protein by Peptide
and Small Molecule Antagonists**



DECLARATION AND POWER OF ATTORNEY

As a below named inventor, I declare that:

My residence, post office address and citizenship are as stated below next to my name; I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural inventors are named below) of the subject matter which is claimed and for which a patent is sought on the invention entitled: **MOLECULAR INTERACTIONS IN HEMATOPOIETIC CELLS** the specification of which X is attached hereto or _____ was filed on _____ as Application No. _____ and was amended on _____ (if applicable).

I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above. I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, Section 1.56. I claim foreign priority benefits under Title 35, United States Code, Section 119 of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)

Country	Application No.	Date of Filing	Priority Claimed Under 35 USC 119

I hereby claim the benefit under Title 35, United States Code § 119(e) of any United States provisional application(s) listed below:

Application No.	Filing Date
60/196,460	April 11, 2000
60/196,528	April 11, 2000
60/196,527	April 11, 2000
60/196,267	April 11, 2000
60/182,296	February 14, 2000
60/176,195	January 14, 2000
60/170,453	December 13, 1999
60/162,498	October 29, 1999
60/160,860	October 21, 1999
60/134,118	May 14, 1999
60/134,117	May 14, 1999
60/134,114	May 14, 1999

I claim the benefit under Title 35, United States Code, Section 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, Section 112, I acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, Section 1.56 which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

Application No.	Date of Filing	Status
Unassigned	October 13, 2000	Pending
09/570,118	May 12, 2000	Pending
09/570,364	May 12, 2000	Pending
09/569,525	May 12, 2000	Pending
09/547,276	April 11, 2000	Pending

POWER OF ATTORNEY: As a named inventor, I hereby appoint the following attorney(s) and/or agent(s) to prosecute this application and transact all business in the Patent and Trademark Office connected therewith.

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I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application or any patent issuing thereon.

Signature of Inventor 1	Signature of Inventor 2	Signature of Inventor 3
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